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<p>(54) Title: POLYKETIDE-ASSOCIATED SUGAR BIOSYNTHESIS GENES</p>		
<p>(57) Abstract</p> <p>The present invention provides isolated polynucleotides from <i>Saccharomyces erythraea</i> that encode enzymes involved in the biosynthesis of polyketide-associated sugars. Methods of using the polynucleotides to produce novel glycosylation modified polyketides are also provided.</p>		

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POLYKETIDE-ASSOCIATED SUGAR BIOSYNTHESIS GENES

This application claims the benefit of U.S. Serial No. 08/576,626 filed December 21, 1995, now pending.

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Field of the Invention

The present invention relates to methods for directing the biosynthesis of specific polyketide analogs by genetic manipulation. In particular, sugar biosynthesis genes are manipulated to produce precise, novel glycosylation-modified macrolides of predicted structure.

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Background of the Invention

Polyketides are a large class of natural products that includes many important antibiotic, antifungal, anticancer, and anti-helminthic compounds such as erythromycins, amphotericins, daunorubicins, and avermectins. Their synthesis proceeds by an ordered condensation of acyl esters to generate carbon chains of varying length, side chain, and reduction pattern that are differentially cyclized and subsequently modified to give the mature polyketides. For many polyketides, maturation includes the addition of one or more sugar residues to the cyclized carbon chain. The sugar residues are frequently critical to the biological activity of the mature polyketide.

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Streptomyces and the closely related *Saccharopolyspora* genera are prodigious producers of polyketide metabolites. Because of the commercial significance of these compounds, a great amount of effort has been expended in the study of *Streptomyces* genetics. Consequently, much is known about *Streptomyces* and several cloning vectors exist for introducing DNA into these organisms.

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Although many polyketides have been identified, there remains the need to obtain novel glycosylation modified (as defined herein) polyketide structures with enhanced properties. Current methods of obtaining such molecules include screening of biological samples and chemical modification of existing polyketides, both of which are costly and time consuming. Current screening methods are based on gross properties of the molecule, i.e. antibacterial, antifungal activity, etc., and both *a priori* knowledge of the structure of the molecules obtained or predetermination of enhanced properties are virtually impossible. Standard chemical modification of existing structures has been successfully employed, but is limited by the number of types of compounds obtainable. Furthermore, the poor yield of multistep chemical syntheses often limits the practicality of this approach. The following modifications to sugar residues bound to polyketides are particularly difficult or inefficient at the present time: change the stereochemistry of specific hydroxyl or methyl groups, change the oxidation state of specific hydroxyl groups, and deoxygenation of specific carbons. Accordingly, there exists a need to obtain molecules wherein such changes are specified and

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performed which would represent an improvement in the technology to produce altered glycosylation-modified polyketide molecules with predicted structure.

The present invention overcomes these problems by providing the genetic sequence of sugar biosynthesis genes involved in the biosynthesis of polyketide-associated sugars.

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Summary of the Invention

In one aspect, the present invention provides an isolated single or double stranded polynucleotide, typically DNA, having a nucleotide sequence which comprises (a) a nucleotide sequence selected from the group consisting of (i) the sense sequence of FIG. 4A (SEQ ID NO:1) from about nucleotide position 54 to about nucleotide position 1136; (ii) the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412; (iii) the sense sequence of SEQ ID NO:1 from about nucleotide position 2409 to about nucleotide position 3410 ; (iv) the sense sequence of FIG. 4B (SEQ ID NO:2) from about nucleotide position 80 to about nucleotide position 1048; (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295; (vi) the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061; (vii) the sense sequence of SEQ ID NO:2 from about nucleotide position 3214 to about nucleotide position 4677; (viii) the sense sequence of SEQ ID NO:2 from about nucleotide position 4674 to about nucleotide position 5879; (ix) the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386; and (x) the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996; (b) sequences complementary to the sequences of (a); (c) sequences that, on expression, encode a polypeptide encoded by the sequences of (a); and (d) analogous sequences that hybridize under stringent conditions to the sequences of (a) and (b). A preferred molecule is a DNA molecule. In another embodiment, the polynucleotide is an RNA molecule.

In another embodiment, a DNA molecule of the present invention is contained in an expression vector. The expression vector preferably further comprises an enhancer-promoter operatively linked to the polynucleotide. In a preferred embodiment, the DNA molecule in the vector is one of the preferred sequences mentioned above. In an especially preferred embodiment, the DNA molecule in the vector is the sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048.

The present invention still further provides for a host cell transformed with a polynucleotide or expression vector of this invention. Preferably, the host cell is a bacterial cell selected from the group consisting of *Saccharopolyspora spp.*, *Streptomyces spp.* and *E. coli*.

The present invention also provides methods to produce novel glycosylation modified

polyketide structures by designing and introducing specified changes in the DNA governing the synthesis and attachment of sugar residues to polyketides. According to one method, the biosynthesis of specific glycosylation-modified polyketides is accomplished by genetic manipulation of a polyketide-producing microorganism comprising the steps of isolating a sugar biosynthesis gene-containing DNA sequence from those described above; identifying within the gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to the polyketide; creating one or more specified changes into the DNA fragment or fragments, thereby resulting in an altered DNA sequence; introducing the altered DNA sequence into a polyketide-producing microorganism to replace the original sequence whereby the altered DNA sequence, when translated, results in altered enzymatic activity capable of effecting the production of the specific glycosylation-modified polyketide; growing a culture of the altered polyketide-producing microorganism under conditions suitable for the formation of the specific glycosylation-modified polyketide; and isolating said specific glycosylation-modified polyketide from the culture.

In a second method the biosynthesis of specific glycosylation-modified polyketides is accomplished by isolating a sugar biosynthesis gene-containing DNA sequence from from those described above; identifying within the gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to the polyketide; reversing the strand orientation of the DNA fragment or fragments, thereby resulting in an altered DNA sequence which, when transcribed, results in production of an antisense mRNA; introducing the altered DNA sequence into a polyketide-producing microorganism having an mRNA capable of binding to the antisense mRNA which results in altered enzymatic activity capable of effecting the production of the specific glycosylation-modified polyketide; growing a culture of the altered polyketide-producing microorganism under conditions suitable for the formation of the specific glycosylation-modified polyketide; and isolating the specific glycosylation-modified polyketide from the culture.

In a third method the biosynthesis of specific glycosylation-modified polyketides is accomplished by isolating a sugar biosynthesis gene-containing DNA sequence from from those described above; identifying within the gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to the polyketide; introducing the DNA fragment or fragments into a polyketide-producing microorganism whereupon transcription and translation of the DNA fragment or fragments generate an altered polyketide-producing microorganism that is capable of producing the specific glycosylation-modified polyketide; growing a culture of the polyketide-producing microorganism containing the DNA fragment or fragments under

conditions suitable for the formation of the specific glycosylation-modified polyketide; and isolating the specific glycosylation-modified polyketide from the culture.

Preferably, the sugar biosynthesis gene-containing DNA sequence of the processes described above comprises genes which encode an enzymatic activity involved in the biosynthesis of L-mycarose and/or D-desosamine. More preferably, the sugar biosynthesis gene-containing DNA sequence comprises the sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048.

The present invention is especially useful in manipulating sugar biosynthesis genes from *Streptomyces* and *Saccharopolyspora*, organisms that provide over one-half of the clinically useful antibiotics.

Brief Description of the Drawings

FIG. 1A illustrates the organization of the erythromycin biosynthetic gene cluster and the genetic designations of the biosynthetic genes; FIG. 1B illustrates an abbreviated erythromycin biosynthetic scheme that broadly associates the biosynthetic genes with their role in erythromycin biosynthesis. Seven *eryB* genes, *eryBI* - *eryBVII*, are responsible for the biosynthesis of L-mycarose or its attachment to the erythronolide B ring, and six *eryC* genes, *eryCI* - *eryCVI*, are responsible for the biosynthesis of D-desosamine or its attachment to 3- α -mycarosylerythronolide B. The dashed arrows indicate that the pathway through erythromycin B is not the principal natural biosynthetic route to erythromycin A.

FIG. 2 illustrates the proposed scheme for the biosynthesis of L-mycarose and the *eryB* genes responsible for the specific steps.

FIG. 3 illustrates the proposed scheme for the biosynthesis of D-desosamine and the *eryC* genes responsible for the specific steps.

FIG. 4A(1-4) illustrates the nucleotide sequence (SEQ ID NO:1) of the sugar biosynthesis genes *eryCII* (coordinates 54-1136), *eryCIII* (coordinates 1147-2412), and *eryBII* (coordinates 2409-3410), with corresponding translation of the open reading frames (SEQ ID NO:3, SEQ ID NO:4 and SEQ ID NO:5 respectively). Standard one letter codes for the amino acids appear beneath their respective nucleic acid codons as described herein.

FIG. 4B(1-9) illustrates the nucleotide sequence (SEQ ID NO:2) of the sugar biosynthesis genes *eryBIV* (coordinates 80-1048), *eryBV* (coordinates 1048-2295), *eryCVI* (coordinates 2348-3061), *eryBVI* (coordinates 3214-4677), *eryCIV* (coordinates 4674-5879), *eryCV* (coordinates 5917-7386), and *eryBVII* (coordinates 7415-7996) with corresponding

translation of the putative open reading frames (SEQ ID NO:6, SEQ ID NO:7, SEQ ID NO:8, SEQ ID NO:9, SEQ ID NO:10, SEQ ID NO:11 and SEQ ID NO:12 respectively). Standard one letter codes for the amino acids appear beneath their respective nucleic acid codons as described herein.

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FIG. 5A illustrates the amino acid sequence identity between the sugar biosynthesis enzyme encoded by the *eryBIV* gene of *Sac. erythraea* (SEQ ID NO:6) and the sugar biosynthesis enzymes encoded by the *ascF* gene of *Yersinia pseudotuberculosis* [Thorson *et al.*, *J. Bacteriol.*, 176:5483 (1994)], (SEQ ID NO:13), the *rfbJ* gene of *Salmonella enterica* [Jiang *et al.*, *Mol. Microbiol.*, 5:695 (1991)]. (SEQ ID NO:14), the *strL* gene of *Streptomyces griseus* [Pissowatzki *et al.*, *Mol. Gen. Genet.* 241:193 (1993)] (SEQ ID NO:15) and the *galE* gene of *Escherichia coli* [Lemaire and Hill, *Nucl. Acids Res.* 14:7705 (1986)] (SEQ ID NO:16). In this and all other Figures in which amino acid sequence identity is compared capitalized letters represent consensus (identical) amino acids between species or amino acids which are conservative substitutions for the consensus residues. Also in each Figure, the sequence identified as "consensus" is merely a convenient representation of conserved amino acids and is not intended as a representation of any existing polypeptide sequence.

FIG. 5B illustrates the amino acid sequence identity between the sugar biosynthesis enzyme encoded by the *eryBVII* gene of *Sac. erythraea* (SEQ ID NO:12) and the sugar biosynthesis enzymes encoded by the *strM* gene of *Streptomyces griseus* [Pissowatzki *et al.*, *Mol. Gen. Genet.* 241:193 (1993)] (SEQ ID NO:17), the *rfbC* gene of *Salmonella enterica* [Jiang *et al.*, *Mol. Microbiol.*, 5:695 (1991)] (SEQ ID NO:18), the *rfbF* gene of *Yersinia enterocolitica* [Zhang *et al.*, *Mol. Microbiol.*, 9:309 (1993)] (SEQ ID NO:19), and the *ascE* gene of *Yersinia pseudotuberculosis* [Thorson *et al.*, *J. Bacteriol.*, 176:5483 (1994)] (SEQ ID NO:20).

FIG. 5C illustrates the amino acid sequence identity between the sugar biosynthesis enzyme encoded by the *eryCIV* gene of *Sac. erythraea* (SEQ ID NO:10) and the sugar biosynthesis enzymes encoded by the *eryCI* gene of *Sac. erythraea* [Dhillon *et al.*, *Mol. Microbiol.*, 3:1405 (1989)] (SEQ ID NO:21), the *ascC* gene of *Yersinia pseudotuberculosis* [Weigel *et al.*, *Biochemistry*, 31:2129 (1992), Thorson *et al.*, *J. Am. Chem. Soc.*, 115:6993 (1993), Thorson *et al.*, *J. Bacteriol.*, 176:5483 (1994)] (SEQ ID NO:22), the *dnrJ* gene of *Streptomyces peucetius* [Stutzman-Engwall *et al.*, *J. Bacteriol.*, 174:144 (1992)] (SEQ ID NO:23), the *prgI* gene of *Streptomyces alboniger* [Lacalle *et al.*, *EMBO J.*, 11:785 (1992)] (SEQ ID NO:24), and the *strS* gene of *Streptomyces griseus* [Distler *et al.*, *Gene*, 115:105 (1992)] (SEQ ID NO:25).

FIG. 5D illustrates the amino acid sequence identity between the sugar biosynthesis enzymes encoded by the *eryBV* and *eryCIII* genes of *Sac. erythraea* (SEQ ID NO:7 and SEQ ID NO:4 respectively) and the sugar biosynthesis enzyme encoded by the *dnrS* gene of

FIG. 5E illustrates the amino acid sequence identity between the sugar biosynthesis enzyme encoded by the *eryCVI* gene of *Sac. erythraea* (SEQ ID NO:8) and the sugar biosynthesis enzymes encoded by the *srnX* gene of *Streptomyces ambofaciens* [Geistlich *et al.*, *Mol. Microbiol.*, 6:2019 (1992)] (SEQ ID NO:27), the *rdmD* gene of *Streptomyces purpurascens* [GenBank Accession: U10405] (SEQ ID NO:28) and the glycine methyltransferase of *Rattus norvegicus* [Ogawa *et al.*, *Eur. J. Biochem.* 168:141 (1987)] (SEQ ID NO:29).

FIG. 6A through 6D illustrate the compounds conceivably formed in Examples 1-4 respectively and are representative of compounds formed from Type I (FIG. 6A), Type II (FIG. 6B), and Type III (FIGS. 6C and 6D) alterations.

FIG. 7 illustrates the construction of the expression plasmid pASX2 described in Example 2. For FIGS 7-13 the following abbreviations have been used: *amp*, ampicillin resistance gene; *tsr*, thiostrepton resistance gene; ROP, repressor of plasmid synthesis gene; *eryBI*, *eryBII*, *eryBIII*, *eryBIV*, *eryBV*, *eryBVI*, *eryBVII*, *eryCI*, *eryCII*, *eryCIII*, *eryCIV*, *eryCV*, and *eryCVI*, the erythromycin biosynthetic genes involved in the synthesis of mycarose or its attachment to the macrolide ring (*eryB*) or the synthesis of desosamine or its attachment to the macrolide ring (*eryC*) [the thin arrows above a gene indicate its relative size and the direction of transcription]; ori-*E. coli*, an origin of DNA replication that functions in *E. coli*, in the specific examples the ColE1 origin; ori-*Streptomyces*, an origin of DNA replication that functions in *Streptomyces*, in the specific examples the pJV1 origin [Servin-Gonzalez *et al.*, *Microbiology*, 141:2499 (1995)]; p-*ermE**, a modified promoter for the erythromycin resistance gene; t-fd, the gene VIII transcription terminator of bacteriophage fd; PCR, polymerase chain reaction. Restriction enzyme sites have been indicated by their standard commercial names (i.e. *Bam*HI, *Eco*RI, etc). The abbreviations appended to the large arrows in the plasmid synthetic schemes summarize each of the steps involved the plasmid constructions. These steps are described fully in the relevant Examples.

FIG. 8 illustrates the construction of the *eryBVII* antisense expression plasmid pASBVII described in Example 2.

FIG. 9A illustrates the construction of the carrier plasmid pK1.

FIG. 9B-E illustrates the construction of plasmid pKB6 which carries all of the *eryB* genes and is described in Example 3.

FIG. 10 illustrates the construction of expression plasmid pX1 described in Example 3.

FIG. 11 illustrates the construction of the *eryB* expression plasmids pXSB6 and pXB6 described in Example 3.

FIG. 12A-B illustrate the construction of plasmid pKC4 which carries all of the *eryC* genes described in Example 4.

FIG. 13 illustrates the construction of the *eryC* expression plasmids pXSC4 and pXC4 described in Example 4.

Detailed Description of the Invention

I. The Invention

The present invention provides isolated and purified polynucleotides that encode enzymes or fragments thereof responsible for the biosynthesis of polyketide-associated sugars or their attachment to polyketides, vectors containing those polynucleotides, host cells transformed with those vectors, a process of making novel glycosylated polyketides using those polynucleotides and vectors, and isolated and purified recombinant polypeptides and polypeptide fragments thereof.

II. Definitions

For the purposes of the present invention as disclosed and claimed herein, the following terms are defined.

The term "polyketide" as used herein refers to a large and diverse class of natural products, including but not limited to antibiotic, antifungal, anticancer, and anti-helminthic compounds. Antibiotics include, but are not limited to anthracyclines and macrolides of different types (polyenes and avermectins as well as classical macrolides such as erythromycins).

The term "glycosylated polyketide" refers to any polyketide that contains one or more sugar residues.

The term "glycosylation-modified polyketide" refers to a polyketide having a changed glycosylation pattern or configuration relative to that particular polyketide's unmodified or native state.

The term "polyketide-producing microorganism" as used herein includes any
 5 microorganism that can produce a polyketide naturally or after being suitably engineered (i.e. genetically). Examples of actinomycetes and the polyketides they naturally produce include but are not limited to those listed in Table 1 below (see Hopwood, D.A. and Sherman, D.H., *Annu. Rev. Genet.*, 24:37-66 (1990) incorporated herein by reference).

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Table 1

Organism	Polyketide Produced
<i>Saccharopolyspora erythraea</i>	Erythromycin
<i>Streptomyces ambofaciens</i>	Spiramycin
<i>Streptomyces avermitilis</i>	Avermectin
<i>Streptomyces fradiae</i>	Tylosin
<i>Streptomyces griseus</i>	Candididin, monactin, griseusin
<i>Streptomyces violaceoniger</i>	Granaticin
<i>Streptomyces thermotolerans</i>	Carbomycin
<i>Streptomyces rimosus</i>	Oxytetracycline
<i>Streptomyces peucetius</i>	Daunorubicin
<i>Streptomyces coelicolor</i>	Actinorhodin
<i>Streptomyces glaucescens</i>	Tetracenomycin
<i>Streptomyces roseofulvus</i>	Frenolicin
<i>Streptomyces cinnamomensis</i>	Monensin
<i>Streptomyces curacoi</i>	Curamycin
<i>Amycolatopsis mediterranei</i>	Rifamycin

Other examples of polyketide-producing microorganisms that produce polyketides naturally include various *Actinomadura*, *Dactylosporangium* and *Nocardia* strains.

The term "sugar biosynthesis genes" as used herein refers to sequences of DNA from
 15 *Saccharopolyspora erythraea* that encode sugar biosynthesis enzymes and is intended to include sequences of DNA from other polyketide-producing microorganisms which are identical or analogous to those obtained from *Saccharopolyspora erythraea*.

The term "sugar biosynthesis enzymes" as used herein refers to polypeptides which are involved in the biosynthesis and/or attachment of polyketide-associated sugars and their
 20 derivatives and intermediates.

The term "polyketide-associated sugar" refers to a sugar that is known to attach to polyketides or that can be attached to polyketides by the processes described herein.

The term "sugar derivative" refers to a sugar which is naturally associated with a polyketide but which is altered relative to the unmodified or native state; examples only include N-3- α -desdimethyl D-desosamine, D-mycarose, 4-keto-L-mycarose, 4-keto-D-mycarose, 3-desmethyl L-mycarose and 3-desmethyl D-mycarose.

The term "sugar intermediate" refers to an intermediate compound produced in a sugar biosynthesis pathway.

The term "*eryB*" as used herein refers to sequences of DNA that encode enzymes involved specifically in the biosynthesis of the deoxysugar L-mycarose.

The term "*eryC*" as used herein refers to sequences of DNA that encode enzymes involved specifically in the biosynthesis of the deoxysugar D-desosamine.

III. Polynucleotides

The organization of the segment of the *Saccharopolyspora erythraea* (*Sac. erythraea*) chromosome that determines the biosynthesis of erythromycin and the corresponding genes that determine the biosynthesis of the sugars L-mycarose and D-desosamine, designated *eryB* and *eryC*, respectively, are shown in FIG. 1A. It is seen that several genes are required for the biosynthesis of each of the sugars and that these genes are interspersed among one another. It is predicted that each gene encodes an enzyme that catalyzes one or a few steps in the biosynthesis of L-mycarose or D-desosamine from thymidine diphospho-4-keto-6-deoxyglucose (TDP-glucose); these steps are outlined in FIG. 2 and FIG. 3. In the case of L-mycarose, (shown in FIG. 2), these steps include: (1) C-2" deoxygenation, (2) C-2"/C-3" enoyl reduction, (3) C-5" epimerization, (4) C-3" C-methylation, (5) C-4" keto reduction, and (6) transfer to erythronolide B. For D-desosamine, shown in FIG. 3, these steps comprise (1) C-4'/3' isomerization, (2, 3) C-3' deoxygenation and reduction, (4) C-3' amination, (5, 6) N-3a' N-dimethylation, and transfer to mycarosyl erythronolide B.

This classification of genes (as belonging to either the *eryB* class or *eryC* class) was determined by first altering the wild type genes of interest in an erythromycin producing strain (i.e. *in vivo*) to inactivate their expression. The erythromycin products resulting from such alterations were then analyzed. Genes whose alterations caused an accumulation of erythronolide B (indicating a lack of L-mycarose, or failure to attach L-mycarose to the erythronolide ring) were classified as *eryB* genes; genes whose alterations caused an accumulation of 3- α -L-mycarosyl erythronolide B (indicating a lack of D-desosamine, or failure to attach D-desosamine to the 3- α -L-mycarosyl erythronolide B ring) were classified as *eryC* genes. Accordingly, it should be noted that all such genes identified herein as *eryB* or *eryC* are involved in the synthesis of L-mycarose or D-desosamine. The predicted

functional activities of the polypeptides encoded by *eryB* and *eryC* will be discussed in further detail below.

In one aspect then, the present invention provides isolated and purified *eryB* and *eryC* polynucleotides from *Sac. erythraea* that encode enzymes involved in the production of glycosylated polyketides. A polynucleotide of the present invention that encodes a sugar biosynthesis enzyme is an isolated single or double stranded polynucleotide having a nucleotide sequence which comprises (a) a nucleotide sequence selected from the group consisting of (i) the sense sequence of FIG. 4A (SEQ ID NO:1) from about nucleotide position 54 to about nucleotide position 1136; (ii) the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412; (iii) the sense sequence of SEQ ID NO:1 from about nucleotide position 2409 to about nucleotide position 3410 ; (iv) the sense sequence of FIG. 4B (SEQ ID NO:2) from about nucleotide position 80 to about nucleotide position 1048; (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295; (vi) the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061; (vii) the sense sequence of SEQ ID NO:2 from about nucleotide position 3214 to about nucleotide position 4677; (viii) the sense sequence of SEQ ID NO:2 from about nucleotide position 4674 to about nucleotide position 5879; (ix) the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386; and (x) the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996;

- (b) sequences complementary to the sequences of (a),
- (c) sequences that, when expressed, encode polypeptides encoded by the sequences of (a), and
- (d) analogous sequences that hybridize under stringent conditions to the sequences of (a).

A preferred polynucleotide is a DNA molecule. In another embodiment, the polynucleotide is an RNA molecule.

The nucleotide sequence and deduced amino acid residue sequences of the sugar biosynthesis genes are set forth in FIG. 4A(1-4) and FIG. 4B(1-9). The nucleotide sequences of FIG. 4A(1-4) (SEQ ID NO:1) and FIG. 4B(1-9) (SEQ ID NO:2) represent full length DNA clones of the sense strand of two distinct clusters of sugar biosynthesis genes and are intended to represent both the sense strand (shown on top) and its complement. The amino acid sequences depicted below the sense strand correspond to polypeptides encoded by a nucleotide sequence selected from the group consisting of (i) the sense strand of SEQ ID NO:1 from about nucleotide position 54 to about nucleotide position 1136 (ii) the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412, (iii) the sense sequence of SEQ ID NO:1 from about nucleotide position 2409 to about

nucleotide position 3410, (iv) the sense sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048, (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295, (vi) the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061, (vii) the sense sequence of SEQ ID NO:2 from about nucleotide position 3214 to about nucleotide position 4677, (ix) the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386 and (x) the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996. The polypeptides encoded by the nucleotide sequences of (i)-(x) above are set forth as SEQ ID NO:3-SEQ ID NO:12 respectively.

The present invention also contemplates analogous DNA sequences which hybridize under stringent hybridization conditions to the DNA sequences set forth above. Stringent hybridization conditions are well known in the art and define a degree of sequence identity greater than about 80%-90%. The modifier "analogous" refers to those nucleotide sequences that encode analogous polypeptides (i.e. in relation to a sugar biosynthesis enzyme), analogous polypeptides being those which have only conservative differences and which retain the conventional characteristics and activities of sugar biosynthesis enzymes. (A more detailed description of analogous polypeptides is provided below). The present invention also contemplates naturally occurring allelic variations and mutations of the DNA sequences set forth above so long as those variations and mutations code, on expression, for a sugar biosynthesis gene of this invention as set forth hereinafter.

As is well known in the art, because of the degeneracy of the genetic code, there are numerous other DNA and RNA molecules that can code for the same polypeptides as those encoded by the aforementioned sugar biosynthesis genes and fragments thereof. The present invention, therefore, contemplates those other DNA and RNA molecules which, on expression, encode the polypeptides of SEQ ID NO:3-SEQ ID NO:11 or fragments thereof. Having identified the amino acid residue sequence encoded by a sugar biosynthesis gene, and with knowledge of all triplet codons for each particular amino acid residue, it is possible to describe all such encoding RNA and DNA sequences. DNA and RNA molecules other than those specifically disclosed herein and, which molecules are characterized simply by a change in a codon for a particular amino acid, are within the scope of this invention.

The 20 common amino acids and their representative abbreviations, symbols and codons are well known in the art (see for example, *Molecular Biology of the Cell*, Second Edition, B. Alberts *et al.*, Garland Publishing Inc., New York and London, 1989). As is also well known in the art, codons constitute triplet sequences of nucleotides in mRNA molecules and as such, are characterized by the base uracil (U) in place of base thymidine (T) which is present in DNA molecules. A simple change in a codon for the same amino acid residue

within a polynucleotide will not change the structure of the encoded polypeptide. By way of example, it can be seen from SEQ ID NO:1 that an AGC codon for serine exists at nucleotide positions 126-128 and again at positions 420-422 and 561-563. However, it can also be seen from that same sequence that serine can be encoded by a TCG codon (see eg. nucleotide
5 positions 192-194) and a TCC codon (see e.g., nucleotide positions 204-206). Substitution of the latter codons for serine with the AGC codon for serine, or *visa versa*, does not substantially alter the DNA sequence of SEQ ID NO:1 and results in production of the same polypeptide. In a similar manner, substitutions of the recited codons with other equivalent codons can be made in a like manner without departing from the scope of the present
10 invention.

A polynucleotide of the present invention can also be an RNA molecule. An RNA molecule contemplated by the present invention is complementary to or hybridizes under stringent conditions to any of the DNA sequences set forth above. Exemplary and preferred RNA molecules are mRNA molecules that encode sugar biosynthesis enzymes of this
15 invention.

IV. Polypeptides

In another aspect, the present invention provides polypeptides which are reasonably believed to be sugar biosynthesis enzymes. A sugar biosynthesis enzyme of the present
20 invention is a polypeptide of about 21 kdal to about 47 kdal. As set forth in FIG. 5A-5E, analogs of the predicted polypeptides encoded by certain *eryB* and *eryC* genes have been identified in various species and their sequences compared using the PRETTY routine (Genetics Computer Group (GCG) Sequence Analysis Software Package, Madison, WI). Due to the degree of amino acid sequence identity existing between the polypeptides of these
25 other sugar biosynthesis genes and the polypeptides encoded by the *eryB* and *eryC* genes, certain enzymatic activities can reasonably be attributed to the *eryB* and *eryC* polypeptides.

By way of example, analogs of the polypeptide encoded by the *eryBIV* gene have been identified in *Yersinia pseudotuberculosis*, *Salmonella enterica*, *Streptomyces griseus* and *Escherichia coli* (see FIG. 5A). The various analogs have been identified with from 290-328
30 amino acid residues and are characterized by a low degree of amino acid sequence identity. (For example, the identity between the sugar biosynthesis enzyme encoded by the *eryBIV* gene of *Sac. erythraea* and the sugar biosynthesis enzyme encoded by the *galE* gene of *E. coli* is 20% at the amino acid level). However, a conserved amino acid sequence motif, G x x G x x G (where G represents the amino acid glycine and x represents any other amino acid
35 residue) is found within the first 30 amino acid residues of all analogs shown. Since the polypeptide encoded by the *galE* gene has been shown to be an epimerase (whose mechanism includes a ketoreduction (Bauer *et al.*, *Proteins* 12:372 (1992)), the *eryBIV* gene product is

reasonably predicted to be a ketoreductase.

As set forth in FIG. 5B analogs of the sugar biosynthesis enzyme encoded by the *eryBVII* gene have been identified in *Streptomyces griseus*, *Salmonella enterica*, *Yersinia enterocolitica* and *Yersinia pseudotuberculosis*. The various analogs have been identified with
5 from 183-200 amino acid residues and are characterized by a moderate degree of amino acid identity. By way of example, the identity at the amino acid level between the sugar biosynthesis enzyme encoded by the *eryBVII* gene of *Sac. erythraea* and the sugar biosynthesis enzyme encoded by the *rfbC* gene of *Salmonella enterica* or the *strM* gene of *Streptomyces griseus* is 37% and 61%, respectively. Furthermore, a common characteristic
10 of these particular polypeptides (including that of *eryBVII*), is that they are only associated with L-sugar biosynthesis and not with D-sugar biosynthesis. Thus the gene product of *eryBVII* is reasonably predicted to function as a C-5 epimerase which converts the stereochemistry of the sugar from the "D" configuration to the "L" configuration.

As set forth in FIG. 5C analogs of the sugar biosynthesis enzyme encoded by the
15 *eryCIV* gene have been identified in *Sac. erythraea* and *Yersinia pseudotuberculosis*. As set forth in FIG. 5C, the predicted amino acid sequences of the protein products of *eryCI* and *eryCIV* share 34% sequence identity to each other, 27% and 25% respectively to the predicted amino acid sequence encoded by *ascC* from *Yersinia pseudotuberculosis*. The enzyme encoded by *ascC* has been shown to remove a hydroxyl group located at the C-3
20 position of L-ascarylose (Liu and Thorson, *Annu. Rev. Microbiol.* 48:223 (1994)). Thus, at least one of the polypeptides encoded by *eryCI* or *eryCIV* is predicted to be an enzyme which functions in deoxygenation reactions.

Furthermore, the enzyme encoded by the *ascC* gene requires the biochemical cofactor pyridoxamine, which is the same cofactor used in biochemical transamination reactions.
25 Consequently, it has been proposed that some protein analogs (such as *dnrJ* from *Streptomyces peucetius*, *prg1* from *Streptomyces alboniger* and *strs* from *Streptomyces griseus*) having a moderate degree of sequence similarity to the polypeptide encoded by *ascC* function as transaminases in amino sugar biosynthesis (Thorson *et al.*, *J. Am. Chem. Soc.* 115:6993 (1993)). Since the biosynthesis of D-desosamine requires both deoxygenation and
30 transamination, it is reasonable to predict that at least one of the polypeptides encoded by the *eryCI* or *eryCIV* genes functions in transamination reactions.

As set forth in FIG. 5D the predicted polypeptides encoded by *eryBV* and *eryCIII* share 43% identity at the amino acid level and as such, may be assumed to have similar activities with respect to their particular sugars. However, as shown in FIGS. 2 and 3, there
35 are no common steps in the proposed pathways of L-mycarose and D-desosamine biosynthesis. Rather than having similar sugar biosynthesis functions, these polypeptides are predicted to be nucleotidyl-sugar transferases which, (in *Sac. erythraea* at least), function to

attach L-mycarose and D-desosamine to erythronolide B and 3- α -mycarosylerythronolide B, respectively.

As set forth in FIG. 5E analogs of the polypeptide encoded by the *eryCVI* gene have been identified in *Streptomyces ambofaciens*, *Streptomyces purpurascens*, and *Rattus norvegicus*. The various analogs have been identified with from 237-293 amino acid residues and are characterized by a low to moderate degree of amino acid identity. By way of example, the identity between the polypeptide encoded by the *eryCVI* gene of *Sac. erythraea* and the glycine methyltransferase of *Rattus norvegicus* is 26% at the amino acid level. Furthermore these sugar biosynthesis enzymes share a common sequence motif, LDVACGTG (SEQ ID NO:30 = amino acid positions 64-71 in the consensus sequence in FIG. 5E), with rat glycine methyltransferase whose biochemical function is known (Ogawa *et al.*, *Eur. J. Biochem.* 168:141 (1987)). Thus these polypeptides are predicted to be N-methyltransferases.

In another aspect, the present invention provides a recombinant C-4" keto reductase from *Sac. erythraea*. A recombinant *Sac. erythraea* C-4" ketoreductase of the present invention is a polypeptide of about 322 or less amino acid residues. A preferred recombinant *Sac. erythraea* C-4" ketoreductase is that encoded by the nucleotide sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048.

The present invention also contemplates amino acid residue sequences that are substantially duplicative of the sequences set forth herein such that those sequences demonstrate like biological activity to disclosed sequences. Such contemplated sequences include those analogous sequences characterized by a minimal change in amino acid residue sequence or type (e.g., conservatively substituted sequences) which insubstantial change does not alter the fundamental nature and biological activity of the aforementioned sugar biosynthesis enzymes.

It is well known in the art that modifications and changes can be made in the structure of a polypeptide without substantially altering the biological function of that peptide. For example, certain amino acids can be substituted for other amino acids in a given polypeptide without any appreciable loss of function. In making such changes, substitutions of like amino acid residues can be made on the basis of relative similarity of side-chain substituents, for example, their size, charge, hydrophobicity, hydrophilicity, and the like.

As detailed in United States Patent No. 4,554,101, incorporated herein by reference, the following hydrophilicity values have been assigned to amino acid residues: Arg (+3.0); Lys (+3.0); Asp (+3.0); Glu (+3.0); Ser (+0.3); Asn (+0.2); Gln (+0.2); Gly (0); Pro (-0.5); Thr (-0.4); Ala (-0.5); His (-0.5); Cys (-1.0); Met (-1.3); Val (-1.5); Leu (-1.8); Ile (-1.8); Tyr (-2.3); Phe (-2.5); and Trp (-3.4). It is understood that an amino acid residue can be substituted for another having a similar hydrophilicity value (e.g., within a value of plus or

minus 2.0) and still obtain a biologically equivalent polypeptide.

In a similar manner, substitutions can be made on the basis of similarity in hydrophathic index. Each amino acid residue has been assigned a hydrophathic index on the basis of its hydrophobicity and charge characteristics. Those hydrophathic index values are:
5 Ile (+4.5); Val (+4.2); Leu (+3.8); Phe (+2.8); Cys (+2.5); Met (+1.9); Ala (+1.8); Gly (-0.4); Thr (-0.7); Ser (-0.8); Trp (-0.9); Tyr (-1.3); Pro (-1.6); His (-3.2); Glu (-3.5); Gln (-3.5); Asp (-3.5); Asn (-3.5); Lys (-3.9); and Arg (-4.5). In making a substitution based on the hydrophathic index, a value of within plus or minus 2.0 is preferred.

10 V. Production of novel glycosylated polyketides

In another aspect, the present invention comprises a general procedure for producing novel polyketide structures *in vivo* by selectively altering, inactivating, or augmenting the genetic information of the organism that naturally produces a related polyketide. That is, in the present invention, novel polyketides of desired structure are produced by manipulation of
15 the *eryB* and/or *eryC* genes followed by their introduction into various polyketide-producing microorganisms. These manipulations result in the formation of "glycosylation-modified" polyketides (i.e. polyketides having an altered glycosylation pattern or configuration relative to their native state). For example, "glycosylation-modified" polyketides are those which have additional sugar groups attached (where none previously existed), different sugars (such
20 as sugar intermediates) attached in place of the natural sugars or lack sugar groups (at positions where sugar groups previously existed).

In the case of Type I and Type II alterations (further described below) glycosylation-modified polyketides may arise through mechanisms which cause either (1) the non-production of the sugar attachment enzyme (i.e. the enzyme involved in attachment of a sugar
25 to the the polyketide structure) or (2) the non-production of a sugar biosynthesis enzyme. In the first instance, the sugar will not be attached to the polyketide since the enzyme which functions to attach the sugar will be lacking. In the second situation, a sugar intermediate from the biosynthesis pathway will be produced (depending on which enzyme is lacking) and attached to the polyketide provided it is recognized as a suitable substrate by the sugar
30 attachment enzyme; alternatively, it will not be recognized and therefore, not attached. In the case of Type III alterations (also described in detail below), glycosylation-modified polyketides arise via attachment of additional or different sugars (i.e. not normally found in a particular polyketide-producing strain) to the polyketide. It should be noted, that these postulated mechanisms are simply provided to enhance understanding of the novel processes
35 described herein; the actual mechanisms by which the Type I, II and III alterations produce glycosylation-modified polyketides is not presently known.

In the first type of alteration (referred to herein as Type I alterations), genetically

altered *eryB* and/or *eryC* genes are introduced into the chromosome of *Sac. erythraea* or another glycosylated polyketide-producing organism that also produces L-mycarose, D-desosamine, or their closely related derivatives such as mycaminose (4-hydroxy D-desosamine). The genetic alteration of an *eryB* and/or *eryC* gene is such that it causes a non-functional enzyme to be synthesized. Once introduced into an appropriate strain, the altered gene replaces its corresponding wild type gene causing the strain to lose the ability to produce a particular enzymatic activity involved in sugar biosynthesis. As a result, a glycosylation-modified polyketide is produced via either of the mechanisms previously described for a Type I alteration.

In a Type I change described herein, a specific mutation in an *eryB* and/or *eryC* gene of the *Sac. erythraea* chromosome is accomplished by a three step process which involves: 1) specifically altering the DNA sequence of a desired sugar biosynthesis gene, 2) subcloning the altered sequence into a suitable vector capable of recombining in the chromosome of an appropriate host and 3) introducing the vector containing the subcloned sequence into the appropriate host so that exchange of the wild type allele with the mutated one will occur. The first step is accomplished using standard recombinant DNA techniques to effect a deletion, base pair conversion or frame-shift in the DNA sequence. The second step, which also employs standard recombinant techniques, involves subcloning the altered sequence into a vector which does not replicate in *Sac. erythraea* or the desired host. In the final step, the vector is introduced into a suitable host, where by the process of gene replacement, the altered allele replaces the wild-type one. All techniques employed in a Type I change are well known to those of ordinary skill in the art.

Example 1 illustrates the process of gene replacement of an *eryB* gene. As Example 1 shows, the *eryB* gene of interest is mutated and along with adjacent upstream and downstream DNA sequences, cloned into a non-replicating *Sac. erythraea* plasmid vector. The vector carrying the mutated allele and adjoining DNA is then introduced into the host strain by the process of protoplast transformation. Transformants are regenerated under selective conditions (i.e. conditions that require expression of a particular plasmid marker) in order to induce recombination of the plasmid into the host cell chromosome. In other words, since the plasmid does not replicate autonomously, it must reside in the chromosome to be maintained in the cell and to express a particular marker under selective conditions. Insertion is achieved when the regenerated cells undergo a single homologous recombination between one of the two DNA segments that flank the mutation on the plasmid and its homologous counterpart in the chromosome. The cells are then grown without selection for the marker which induces plasmid loss from the chromosome. This loss arises after the cells have undergone a second recombination between the second DNA segment that flanks the mutation and its homologous chromosomal counterpart. This second recombinational event

results in the loss of the plasmid sequences and the wild type allele from the chromosome; the mutant allele however is retained.

In a variation of a Type I change, the non-production of the sugar biosynthesis enzyme (or attachment enzyme) may be achieved by the alternative mechanisms of promoter inactivation and/or transcriptional terminator insertion. These variations do not effect the gene sequence itself but rather regulatory mechanisms involved in gene transcription. "Promoter" as used herein refers to that region of a DNA molecule which controls the initiation of RNA transcription. Such regions are known to bind RNA polymerases (i.e. the enzymes involved in synthesizing RNA molecules). This form of Type I change (i.e. promoter inactivation) involves two steps of 1) identifying the promoter region of the desired gene and 2) rendering the promoter region inoperable by mutation. As in the replacement mechanism described above such mutations may be effected by creating deletions in the promoter sequence or by base pair conversion. In the case where the promoter controls transcription of a single gene, inactivation of the promoter will eliminate expression of that particular gene; of course, where the promoter controls expression of an entire operon (i.e. a series of genes whose expression is controlled by a single promoter), promoter inactivation will effectively eliminate expression of all genes in that operon.

In a similar manner, the non-production of a sugar biosynthesis enzyme (or attachment enzyme) may arise from inserting a transcriptional terminator upstream from the gene to be inactivated. A "transcriptional terminator" as used herein is a nucleotide sequence which signals RNA polymerase to cease transcription. An example of a transcriptional terminator is a palindromic sequence capable of forming a stem-loop structure that is followed by a stretch of U residues (for example the transcriptional terminator that follows gene VIII of bacteriophage fd (Beck and Zink, *Gene*, 16:35 (1981))). Effecting a change in production of a sugar biosynthesis gene by this process involves 1) identifying of the gene or genes of interest (in the case of an operon arrangement) to be inactivated and 2) cloning a transcriptional terminator sequence in a region of the DNA upstream from such gene(s). A transcriptional terminator will cause the polymerase involved in RNA transcription to stop (at or near the signaling region) thereby preventing transcription of any downstream sequences. Thus, changes such as promoter inactivation and transcriptional insertion, which directly effect expression of sugar biosynthesis genes are also intended to be within the scope of the invention.

In the second case (referred to herein as Type II alterations) *eryB* and/or *eryC* genes are arranged on a vector in an antisense orientation relative to a promoter capable of allowing expression of the gene in *Sac. erythraea* or *Streptomyces*. The vector is then introduced into a polyketide producing microorganism. As a result of this vector construction, antisense messenger RNA (mRNA) is produced which interferes with the translation of the wild-type

mRNA. Similarly to the Type I manipulation, novel glycosylation modified polyketides will be produced in which the normal mycarose, desosamine, and/or closely related sugar residue is lacking or is substituted by a sugar intermediate.

In a Type II change, inactivation of the *eryB* and/or *eryC* genes by antisense expression is accomplished by a two step procedure in which (1) a specific sugar biosynthesis gene is subcloned into an expression vector in an antisense (i.e. reverse) orientation; and (2) the anti-sense expression vector is introduced into the desired strain. The first step is accomplished using standard recombinant DNA techniques employing either *E. coli* or *Streptomyces* as the host, and an expression vector (capable of replicating in either host) that can be assembled to contain a *Streptomyces* promoter. *Streptomyces* promoters may be obtained from any commercially available *Streptomyces* plasmids or *Streptomyces*- *E. coli* shuttle plasmids. In step 2, the anti-sense expression vector is introduced into a suitable *Streptomyces* strain and the transformed cells are grown under selective conditions in order to maintain the expression plasmid in the cell.

As described in Example 2, the gene to be inactivated is subcloned in its reverse orientation downstream of a *Streptomyces* promoter (which is contained within a replicating *Sac. erythraea* plasmid). The plasmid carrying the antisense gene is then introduced into the host strain by protoplast transformation. Transformants are regenerated under selective conditions in order to maintain the autonomously replicating plasmid in the cells. Subsequent expression of the antisense gene causes the production of an antisense messenger RNA (mRNA) that is complementary to the mRNA of the native allele of the selected gene. Through standard nucleotide base pair interactions, the antisense mRNA and the native mRNA form an RNA duplex that occludes the ribosome binding site of the native mRNA. This interaction prevents ribosomal translation of the native mRNA and the corresponding synthesis of the enzyme encoded by that mRNA. In this way, specific enzymatic steps in sugar biosynthesis corresponding to the identity of the gene expressed in the antisense orientation are blocked leading to the production of novel sugar intermediates which, when attached to the polyketide ring of the host microorganism, give rise to novel glycosylation-modified polyketides. Alternatively, the antisense expression vector can be constructed using a non-replicating *Sac. erythraea* vector that includes flanking DNA from a nonessential region of the *Sac. erythraea* chromosome, such as the region immediately upstream from the *eryK* gene (FIG. 1). This vector can then be used to stably insert the antisense construction into the chromosome by homologous recombination in a fashion similar to that described for the construction of a Type I alteration.

In the third case (referred to herein as Type III alterations), novel glycosylation-modified polyketides of desired structure are produced by arranging all or a subset of the *eryB* and/or *eryC* genes on a replicating vector and introducing these genes *en bloc* into a

“distinct” polyketide-producing organism, ie. one other than the microorganism from which the *eryB* and/or *eryC* genes were taken. As an example, *eryB* and/or *eryC* genes may be taken from *Sac. erythraea* and introduced into *Streptomyces violaceoniger* or *Streptomyces venezuelae*. In this case, mycarose, desosamine, their biochemical intermediates and/or their
5 closely related derivatives will be synthesized and attached at specific positions to polyketide compounds that do not necessarily carry these, or any, sugar residues. Some examples of novel glycosylated polyketides that may be produced in hosts that carry such manipulations are shown in FIG. 6.

In Type III changes, the genes for the biosynthesis of mycarose and/or desosamine are
10 introduced into a polyketide-producing organism other than *Sac. erythraea* by another simple two step procedure: 1) all or a subset of the *eryB* and/or *eryC* genes are assembled together on a replicating plasmid downstream of a *Streptomyces* promoter; and 2) the plasmid is introduced into the polyketide-producing organism. Step 1 requires standard recombinant DNA manipulations employing *E. coli* and/or *Streptomyces* as the host. Step 2 requires one
15 or more plasmids out of the several *Streptomyces* vectors or *E. coli-Streptomyces* shuttle vectors available, one or more promoters that function in *Streptomyces*, and a selection for the presence of the strain carrying the plasmid. As described in Examples 3 and 4, sets of the *eryB* and/or *eryC* genes are sequentially subcloned together on a replicating vector downstream of a suitable promoter that functions in the desired host. The plasmid carrying
20 the grouped genes is then introduced into the host strain by electroporation or by transformation of protoplasts employing selection for a plasmid marker.

GENERAL METHODS

25 Materials, Plasmids, and Bacterial Strains

Restriction endonucleases, T4 DNA ligase, competent *E. coli* DH5 α cells, X-gal, IPTG and plasmids pUC18, pUC19, and pBR322 were purchased from Bethesda Research Laboratories (BRL), Gaithersburg, MD. Ventr[®] DNA polymerase was purchased from New
30 England Biolabs (Beverly, MA). Plasmids pGEM[®]5Zf, pGEM[®]7Zf, and pGEM[®]11Zf were from Promega, Madison, WI, plasmids pIJ4070 and pIJ702 were obtained from the John Innes Institute, Norwich, England, and plasmids pWHM3 and pWHM4 (*J. Bacteriol.* 1989 171:5872) were obtained from C. R. Hutchinson, University of Wisconsin, Madison, WI. [α -³²P]dCTP, Hybond[™]-N nylon membranes, and Megaprime nick translation kits were
35 from Amersham Corp., Chicago, IL. SeaKem[®] LE agarose and SeaPlaque[®] low gelling temperature agarose were from FMC Bioproducts, Rockland, ME. *E. coli* K12 strains carrying the *E. coli-Sac. erythraea* shuttle plasmids pWHM3 and pWHM4 (Vara *et al.*, *J*

Bacteriol, 171:5872 (1989)) and pAIX have been deposited at the Agricultural Research Culture Collection (NRRL) 1815 N. University Street, Peoria, Illinois 61604, as of December 5, 1995, under the terms of the Budapest Treaty and will be maintained for a period of thirty (30) years from the date of deposit, or for five (5) years after the last request for the deposit, or for the enforceable period of the U.S. patent, whichever is longer. Plasmids pWHM3, pWHM4 and pAIX were accorded the accession numbers NRRL B-21512, NRRL B-21513 and NRRL B-21514, respectively. *Sac. erythraea* strain NRRL2338 is also available from the Agricultural Research Service culture collection. *Staphylococcus aureus* Th^R (thiostrepton resistant) was obtained by plating 10⁸ cells of *S. aureus* on agar medium containing 10 µg/ml thiostrepton and picking a survivor after 48 hr growth at 37°C. Thiostrepton was obtained from Sigma Chemical, St. Louis, MO. All other chemicals and reagents were from standard commercial sources unless otherwise specified.

DNA Manipulations

Standard conditions were employed for restriction endonuclease digestion, agarose gel-electrophoresis, isolation of DNA fragments from low melting agarose gels, DNA ligation, plasmid isolation from *E. coli* by alkaline lysis, and transformation of *E. coli* employing selection for ampicillin resistance (150 µg/ml) on LB agar plates (Sambrook *et al.*, *Molecular Cloning, A Laboratory Manual*, Cold Spring Harbor Laboratory Press, Plainview, NY, 1989). Total DNA from *Sac. erythraea* and *Streptomyces* species (including *S. fradiae*, *S. celestes*, *S. violaceoniger*, *S. hygroscopicus*, *S. venezuelae*) was prepared according to described procedures (Hopwood *et al.*, *Genetic Manipulation of Streptomyces, A Laboratory Manual*, John Innes Foundation, Norwich, UK (1985)). Transfer of DNA from agarose gels to HybondTM-N membranes and Southern analysis using MegaprimeTM nick translated probes was performed according to the manufacturers instructions.

Amplification of DNA Fragments

Synthetic deoxyoligonucleotides were synthesized on an ABI Model 380A synthesizer (Applied Biosystems, Foster City, CA) following the manufacturers recommendations. Amplification of DNA fragments was performed by the polymerase chain reaction (PCR) using a Perkin Elmer GeneAmp[®] PCR System 9600. Reactions contained 100 pmol of each primer, 1 µg of template DNA (chromosomal DNA from *Sac. erythraea* NRRL2338), 2 units Ventr[®] DNA polymerase in 100 µl volume of PCR buffer (10 mM KCl, 10 mM (NH₄)₂SO₄, 20 mM Tris-HCl (pH 8.8, @ 25°C), 2.5 mM MgSO₄, 0.1% Triton[®] X-100) containing dATP (200 µM), dTTP (200 µM), dCTP (250 µM), and dGTP (250 µM). The reaction mixture was subjected to 30 cycles. Each cycle consisted of one period of 35 sec at 96°C and one period of 2 min at 72°C. The reaction products were visualized and

purified from low melting agarose. The PCR primers described in the examples were derived from the nucleotide sequence of the *eryB* and *eryC* genes of FIG. 4.

Transformation and Gene Replacement in *Sac. erythraea*

5 Protoplasts of *Sac. erythraea* strains were prepared and transformed with miniprep DNA isolated from *E. coli* according to published procedures (Yamamoto *et al.*, *J Antibiotics*, 39:1304 (1986)). Non-integrative transformants, in the case of pWHM4 derivatives, were selected by regenerating the protoplasts and overlaying with thiostrepton (final concentration 20 µg/ml) as described (Weber *et al.*, *Gene*, 68:173 (1988)). Integrative
10 transformants, in the case of pWHM3 derivatives, were selected on thiostrepton-containing agar plates (15 µg/ml) as described by Weber *et al.*, *Gene*, 68:173 (1988). Loss of the Th^R phenotype was monitored after two rounds of non-selective growth in SGGP media (Yamamoto *et al.*, *J Antibiotics*, 39:1304 (1986)) followed by protoplasting and serial
15 dilution on non-selective agar media. Regenerated protoplasts were replica plated on thiostrepton-containing media. Th^S (thiostrepton-sensitive) colonies arose at a frequency of 10⁻¹. Retention of the mutant allele was established by Southern hybridization of several Th^S colonies.

Fermentation

20 *Sac. erythraea* or *Streptomyces* cells are inoculated into 100 ml SCM medium (1.5% soluble starch, 2.0% Difco Soytone, 0.15% Yeast Extract, 0.01% CaCl₂) and allowed to grow for 3 to 6 days. The entire culture is then inoculated into 10 liters of fresh SCM medium. The fermenter is operated for a period of 4 to 7 days at 32°C maintaining constant aeration and pH at 7.0. After the fermentation is complete, the cells are removed by centrifugation at
25 4°C and the fermentation beer is kept cold until further use. When antibiotic selection to maintain a plasmid, such as pXC4 or pXB6, is required, thiostrepton (10µg/ml) is added to both the 100 ml starter culture and the 10-liter fermenter.

30 The invention will be better understood in connection with the following examples, which are intended as an illustration of and not a limitation upon the scope of the invention. Both below and throughout the specification, it is intended that citations to the literature be expressly incorporated by reference.

Example 1: Construction and characterization of *Sac. erythraea* ERBIV that produces 4"-deoxy-4"-oxo-erythromycin A

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A. Construction of Plasmid pRBIV: A 4.3 kb *Pst*I-*Hind*III fragment, which included

the *eryBIV* gene, was isolated from the plasmid pAIX5 and subcloned into *Pst*I-*Hind*III digested pUC19 to generate plasmid pUCBIV. After transformation and isolation of the plasmid from *E. coli*, the identity of pUCBIV was confirmed by digestion with *Mun*I which released a fragment of 370 bp. Plasmid pUCBIV was then cut with the restriction enzyme
5 *Nco*I, the restriction site filled in with Klenow enzyme, and the plasmid religated to generate plasmid pNCOBIV, (which now carried a frameshift mutation in the *eryBIV* gene). After transformation and isolation of the plasmid from *E. coli*, the identity of pNCOBIV was confirmed by digestion with *Nsi*I and *Hind*III which released a fragment of 1.59 kb. (The *Nsi*I site was formed by the fill-in and religation of the *Nco*I site.) Finally, plasmid
10 pNCOBIV was digested with *Hind*III and *Sst*I and the 3.2 kb fragment carrying the altered *eryBIV* gene was isolated and ligated into *Hind*III and *Sst*I digested pWHM3 to generate plasmid pRBIV. After transformation and isolation of the plasmid from *E. coli*, the identity of pRBIV was confirmed by digestion with *Kpn*I which released fragments of 5.2 kb, 4.4 kb, and 0.72 kb.

15 B. Construction of *Sac. erythraea* ERBIV: *Sac. erythraea* protoplasts were transformed with plasmid pRBIV and integrative transformants selected as described in General Methods. Resolution of the integrants by nonselective growth as described in General Methods yielded *Sac. erythraea* ERBIV in which the wild type copy of the *eryBIV* gene was replaced with the inactive mutant copy. Gene replacement was confirmed by
20 Southern analysis of *Nco*I digested *Sac. erythraea* DNA and *Nco*I-*Nsi*I digested *Sac. erythraea* DNA using the 1.58 kb *Nco*I-*Hind*III fragment isolated from plasmid pUCBIV (coordinates 681-2214, FIG. 4B) as a probe. Wild type *Sac. erythraea* and wild type resolvants display a hybridizing DNA fragment of 2.75 kb when digested with either *Nco*I or *Nco*I-*Nsi*I, whereas *Sac. erythraea* strain ERBIV is characterized by hybridization to either a
25 16 kb DNA fragment or a 2.75 kb DNA fragment when digested with *Nco*I or *Nco*I-*Nsi*I, respectively.

C. Isolation, purification, and properties of 4"-deoxy-4"-oxo-erythromycin A from *Sac. erythraea* ERBIV: *Sac. erythraea* strain ERBIV is fermented for 4 days in SCM media as described in General Methods. The fermentation broth of *Sac. erythraea* ERBIV is then
30 cooled to 4°C and adjusted to pH 4.0 and extracted once with methylene chloride. The aqueous layer is readjusted to pH 9.0 and extracted twice with methylene chloride and the combined basic methylene chloride extracts are concentrated to a solid residue. This is digested in methanol and chromatographed over a column of Sephadex LH-20 in methanol. Fractions are tested for bioactivity against a sensitive organism, such as *Staphylococcus aureus* Th^R, and active fractions are combined. The combined fractions are concentrated and
35 the residue is digested in 10 ml of the upper phase of a solvent system consisting of n-heptane, benzene, acetone, isopropanol, 0.05 M, pH 7.0 aqueous phosphate buffer

(5:10:3:2:5, v/v/v/v/v), and chromatographed on an Ito Coil Planet Centrifuge in the same system. Active fractions are combined, concentrated and partitioned between methylene chloride and dilute ammonium hydroxide (pH 9.0). The methylene chloride layer is separated and concentrated to yield the desired product as a white foam.

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Example 2: Construction and characterization of *Sac. erythraea* ER720(pASBVII) that produces 3- α -D-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B

A. Construction of plasmid pASX2 (see FIG. 7): The 290 bp *EcoRI*-*Bam*HI segment carrying the *ermE** promoter is isolated from plasmid pIJ4070 and ligated into *EcoRI*-*Bam*HI digested pWHM4 DNA to form pASX1. After transformation and isolation of the plasmid from *E. coli*, the identity of pASX1 is confirmed by digestion with *Apa*LI which releases fragments of 3.9 kb, 2.5 kb, 1.2 kb, 0.5 kb, and 0.4 kb. Two oligonucleotides of the sequences: SEQ ID NO:31 (5'-GATCCAGCGTCTGCAGGCATGCTCTAGATACAATTA
15 AAGGCTCCTTTTGGAGCCTTTTTTTTTTGGAGATTTTCAACGT-3') and
SEQ ID NO:32 (5'-AGCTACGTTGAAAATCTCCAAAAAAAAAAGGCTCCAAAA
GGAGCCTTTAATTGTATCTAGAGCATGCCTGCAGACGCTG-3'), corresponding to the
(+) and (-) strands of the bacteriophage fd gene VIII transcription terminator (t-fd) (Beck *et al.* (1978) *Nucl. Acids Res.* 5:4495)) and including restriction enzyme sites for the enzymes
20 *Pst*I, *Sph*I, and *Xba*I, and overhanging ends compatible with *Bam*HI and *Hind*III are
synthesized and approximately 250 ng of each oligonucleotide are then mixed together in TE buffer and heated to 99°C for 1 min. The solution is cooled slowly to room temperature allowing the oligonucleotides to anneal due to self complementarity, and the annealed oligonucleotides are then ligated into *Bam*HI-*Hind*III digested pASX1 to give pASX2. After
25 transformation and isolation of the plasmid from *E. coli*, the identity of pASX2 is confirmed by DNA sequencing of the 1.2 kb *EcoRI*-*Sa*II fragment that contains the *ErmE** promoter and the bacteriophage fd terminator.

B. Construction of plasmid pASBVII (see FIG. 8): The 598 base pair DNA segment that carries the *eryBVII* gene, comprising coordinates 7398-7996 (FIG. 4B), is amplified by
30 PCR employing two oligonucleotides, SEQ ID NO:33 (5'-
GATCGCATGCTCTAGAGTACG-TGAGCTGGCGGTGGCGGGC-3') and SEQ ID NO:34
(5'-GATCCGGATCCGCATGCTT-CACCTGCCGGTGCTGGCGGG-3'). After digestion of the purified PCR product with *Bam*HI-*Xba*I the PCR fragment was ligated to *Bam*HI-*Xba*I
digested pASX2 to give pASBVII. After transformation and isolation of the plasmid from *E.*
35 *coli*, the identity of pASBVII is verified by DNA sequencing of the 880 bp *EcoRI*-*Xba*I insert.

C. Construction of *Sac. erythraea* ER720(pASBVII): *Sac. erythraea* strain ER720

protoplasts are transformed with plasmid pASBVII and transformants are selected for with thiostrepton (15 µg/ml). To confirm transformation, total DNA is isolated from Th^R colonies and used to transform *E. coli*. After transformation and isolation of the plasmid from *E. coli*, the identity of pASBVII is verified by restriction analysis with the enzymes *Pvu*II and *Bam*HI which releases a 1.48 kb fragment. Those *Sac. erythraea* colonies that are found to contain pASBVII are designated *Sac. erythraea* ER720(pASBVII).

D. Isolation, purification, and properties of 3- α -D-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B from *Sac. erythraea* ER720(pASBVII): *Sac. erythraea* ER720(pASBVII) is fermented for 3 days in SCM media with thiostrepton selection as described in General Methods. The fermentation broth is then cooled to 4°C and adjusted to pH 4.0 and extracted once with methylene chloride. The aqueous layer is readjusted to pH 9.0 and extracted twice with methylene chloride and the combined extracts are concentrated to a solid residue. This is digested in methanol and chromatographed over a column of Sephadex LH-20 in methanol. Fractions are tested for bioactivity against a sensitive organism, such as *Staphylococcus aureus* Th^R, and active fractions are combined. The combined fractions are concentrated and the residue is digested in 10 ml of the upper phase of a solvent system consisting of n-heptane, benzene, acetone, isopropanol, 0.05 M, pH 7.0 aqueous phosphate buffer (5:10:3:2:5, v/v/v/v/v), and chromatographed on an Ito Coil Planet Centrifuge in the same system. Active fractions are combined, concentrated and partitioned between methylene chloride and dilute ammonium hydroxide (pH 9.0). The methylene chloride layer is separated and concentrated to yield the desired product as a white foam.

Example 3: Construction and characterization of *Streptomyces antibioticus* ATCC 11891(pXB6) that produces 3-des-oleandrosyl-3-mycarosyl oleandomycin

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A. Construction of plasmid pKB6 and intermediates (see FIG. 9)

i) Construction of plasmid pK1: The DNA sequences of pBR322 (GenBank Accession #: J01749) and pUC19 (GenBank Accession #: X02514) are known. The 805 nt DNA segment comprising coordinates 1673 through 2478 of pBR322 is amplified by PCR employing two oligodeoxynucleotides, SEQ ID NO:35 (5'-GATCACATGTTCTTTCCTG-CGTTATCCCCTG-3') and SEQ ID NO:36 (5'-GATCGGATCCATGCATGTCTAGAGCA-TCGCAGGATGCTGCTGGC-3'). After digestion of the purified PCR product with *A*fIII and *Bam*HI the fragment is ligated into *A*fIII and *Bam*HI digested pUC19 to give plasmid pK1. The identity of plasmid pK1, after transformation and isolation from *E. coli*, is verified by *Pvu*II digestion which releases fragments of 0.55 kb and 2.55 kb. Plasmid pK1 contains the ROP region of pBR322 that controls plasmid copy number.

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ii) Construction of plasmid pKB1: The 2.24 kb DNA segment that carries the

eryBIV and *eryBV* genes, comprised between coordinates 56 and 2296 of the sequence presented in SEQ ID NO:2, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:37 (5'-GAATGCATCCTGGAAAGCGAGCAAATGCTCCGGTG-3') and SEQ ID NO:38 (5'-GATCTAGAGCTAGCCGGCGTGGCGGCGCGTG-3'). After digestion with
 5 *NsiI* and *XbaI* the fragment is ligated into *NsiI* and *XbaI* digested pK1 to yield plasmid pKB1, 5.3 kb in size. The identity of plasmid pKB1, after transformation and isolation from *E. coli*, is verified by *KpnI* digestion which releases fragments of 0.72 kb, 1.14 kb and 3.42 kb.

iii) Construction of plasmid pKB2: The 1.56 kb DNA segment that carries the *eryBVI* gene, comprised between coordinates 3121 and 4677 of the sequence presented in
 10 SEQ ID NO:2, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:39 (5'-GATCGCTAGCCGTGACCGGACCCTTACAGTGAGTG-3') and SEQ ID NO:40 (5'-GATCTAGACTTAAGTCATCCGGCGGTCCTGGTGTAGACGGC-3'). After digestion with *NheI* and *XbaI* the fragment is ligated into *NheI* and *XbaI* digested pKB1 to give plasmid pKB2, 6.9 kb in size. The identity of plasmid pKB2, after transformation and isolation from
 15 *E. coli*, is confirmed by *BamHI* digestion which releases fragments of 0.22 kb, 0.40 kb, 2.6 kb and 3.7 kb.

iv) Construction of plasmid pKB3: The 0.6 kb DNA segment that carries the *ery BVII* gene, comprised between coordinates 7385 and 7987 of the sequence presented in
 20 SEQ ID NO:2, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:41 (5'-GATCTTAAGAACCGGAGTTGCGAGTACGTGAGCTGGCG-3') and SEQ ID NO:42 (5'-GATCTAGACCTAGGTCACCTGCCGGTGCTGGCGGGCTC-3'). After digestion with *AflIII* and *XbaI* the fragment is ligated into *AflIII* and *XbaI* digested pKB2 giving plasmid pKB3, 7.5 kb in size. The identity of plasmid pKB3, after transformation and isolation from *E. coli*, is verified by *PstI* digestion which releases fragments of 1.1 kb and 6.4 kb.

25 v) Construction of plasmid pKB4: The 1.0 kb DNA segment that carries the *eryBII* gene, comprised between coordinates 2385 and 3410 of the sequence presented in SEQ ID NO:1, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:43 (5'-GATCCTAGGCCGAGGAAGGAGAGAACCACG-3') and SEQ ID NO:44 (5'-GATCTAGATTAATCACTGCAACCAGGCTTCCGGC-3'). Following digestion with
 30 *AvrII* and *XbaI* the fragment is ligated into *AvrII* and *XbaI* digested pKB3 yielding the desired plasmid pKB4. After transformation and isolation of the plasmid from *E. coli*, the identity of pKB4, 8.5 kb in size, is verified by *BglIII* and *EcoRI* digestion which releases fragments of 0.41 kb, 1.6 kb, 3.1 kb and 3.4 kb.

vi) Construction of plasmid pKB5: The DNA sequence of *eryBIII* has been
 35 reported (Haydock *et al* (1991) *Mol Gen Genet* 230:120). The 1.3 kb DNA segment that carries the *eryBIII* gene, comprised between coordinates 3965 and 5232 of the sequence depicted in Haydock *et al*, is amplified by PCR employing two deoxyoligonucleotides, SEQ

ID NO:45 (5'-GATTAATTGGCCGCGGCGCCGCGCTC-GTTATG-3') and SEQ ID NO:46 (5'-GATCTAGATAATTAATCATACGACTTCCAGTC-GGGGTAG-3'). After digestion with *MseI* and *XbaI* the fragment is ligated into *MseI* and *XbaI* digested pKB4 to give the desired plasmid pKB5, 9.8 kb in size. The identity of pKB5, after transformation and isolation from *E. coli*, is verified by *PstI* digestion which releases fragments of 1.1 kb, 2.5 kb, and 6.1 kb, visualized by gel electrophoresis.

vii) Construction of plasmid pKB6: The *eryBI* gene has been mapped (Haydock *et al* (1991) *Mol Gen Genet* 230:120) and the DNA sequence on both flanks of *eryBI* is known (Haydock *et al* (1991) *Mol Gen Genet* 230:120) and GenBank Accession # M11200. The 2.5 kb DNA segment that carries the *eryBI* gene, comprised between coordinates 1.1 and 3.6 of the map presented in Haydock *et al.*, is amplified by PCR employing two deoxyoligonucleotides: SEQ ID NO:47 (5'-GATTAATTAATGATCA-AGCTGAAAATTGTTTGCATG-3') and SEQ ID NO:48 (5'-GATCTAGACTGCCGGCT-CAGCCTTCCCAGGTTCG-3'). After digestion with *PacI* and *XbaI* the fragment is ligated into *PacI* and *XbaI* digested pKB5 to give plasmid pKB6, 12.3 kb in size. The identity of pKB6, after transformation and isolation from *E. coli*, is verified by *BamHI* digestion which releases fragments of 0.22 kb, 0.40 kb, 1.4 kb, 2.6 kb, 3.3 kb and 4.4 kb. Plasmid pKB6 carries all of the *eryB* genes, *eryBI-eryBVII*, that are involved in the biosynthesis of mycarose and its attachment to the polyketide.

B. Construction of Plasmid pXSB6 (see FIG. 11): The 9.2 kb *NsiI-XbaI* segment of pKB6, prepared as described in Example 3(A)(vii) above, that carries all of the *eryB* genes is isolated and ligated into *PstI-XbaI* digested pASX2, prepared as described in Example 2(A) above, to give plasmid pXSB6. After transformation and isolation of the plasmid from *E. coli*, the identity of pXSB6, 17.2 kb in size, is verified by the observation of fragments of 0.41 kb, 1.9 kb, and 14.9 kb after *EcoRI* digestion. Plasmid pXSB6 carries all of the *eryB* genes in a transcriptional fusion downstream of the *ermE** promoter on an *E. coli-Streptomyces* shuttle plasmid.

C. Construction of Plasmid pXB6

i) Construction of plasmid pN702 (see FIG. 10): Two oligonucleotides of the sequences: SEQ ID NO:49 5'-GGAATTCAGATCTATGCATTCTAGAA-3') and SEQ ID NO:50 (5'-CGCGTTCTAGAATGCATAGATCTGAATTCCTGCA-3') that include restriction enzyme sites for the enzymes *EcoRI*, *BglIII*, *NsiI*, and *XbaI* and overhanging ends compatible with *PstI* and *MluI* are synthesized. Approximately 250 ng of each oligonucleotide are then mixed together in TE buffer and heated to 99°C for 1 min. After the solution is cooled slowly to room temperature allowing the oligonucleotides to anneal due to self complementarity, the annealed oligonucleotides are ligated into *PstI-MluI* digested pIJ702 to yield the desired plasmid pN702. After transformation and isolation of the plasmid

from *Streptomyces lividans* 1326, the identity of plasmid pN702, 4.3 kb in size, is verified by the observation of fragments of 0.75 kb and 3.6 kb after *EcoRI-BamHI* or *XbaI-BamHI* digestion.

5 ii) Construction of plasmid pX1 (see FIG. 10): The 290 bp *EcoRI-BamHI* segment that carries the *ermE** promoter is isolated from plasmid pIJ4070 and ligated into *EcoRI-BglII* digested pN702 to give plasmid pX1. The resulting mixture contains the desired plasmid pX1. After transformation and isolation of the plasmid from *Streptomyces lividans* 1326, the identity of plasmid pX1, 4.6 kb in size, is verified by the observation of fragments of 1.0 kb and 3.6 kb after *NsiI-BamHI* digestion.

10 iii) Construction of plasmid pXB6 (see FIG. 11): The 9.2 kb *NsiI-XbaI* segment of pKB6, prepared as described in Example 3(A)(vii) above, that carries all of the *eryB* genes is isolated and ligated into *NsiI-XbaI* digested pX1 to give the desired plasmid pXB6. After transformation and isolation of the plasmid from *Streptomyces lividans* 1326, the identity of plasmid pXB6, 13.8 kb in size, is verified by the observation of fragments of
15 0.41 kb, 1.9 kb, and 11.5 kb after *EcoRI* digestion. Plasmid pXB6 carries all of the *eryB* genes in a transcriptional fusion to the *ermE** promoter on a *Streptomyces* plasmid.

 D. Construction of *Streptomyces antibioticus* ATCC 11891(pXB6): Approximately 500 µg of plasmid pXB6, isolated from *Streptomyces lividans* 1326(pXB6), are electroporated into the oleandomycin producer *Streptomyces antibioticus* ATCC 11891 and
20 several of the resulting Thio^R colonies that appear on the R3M-agar plates containing thiostrepton are analyzed for their plasmid content. The presence of plasmid pXB6, 13.8 kb in size, is verified by the observation of fragments of 0.41 kb, 1.9 kb, and 11.5 kb after *EcoRI* digestion.

 E. Isolation, purification, and properties of 3-des-oleandrosyl-3-mycarosyl
25 oleandomycin from *Streptomyces antibioticus* ATCC 11891(pXB6): *Streptomyces antibioticus* ATCC 11891(pXB6) is fermented for 5 days in SCM media with thiostrepton selection as described in General Methods. The fermentation broth is then cooled to 4°C and adjusted to pH 4.0 and extracted once with methylene chloride. The aqueous layer is readjusted to pH 9.0 and extracted twice with methylene chloride and the combined extracts
30 are concentrated to a solid residue. This is digested in methanol and chromatographed over a column of Sephadex LH-20 in methanol. Fractions are tested for bioactivity against a sensitive organism, such as *Staphylococcus aureus* Th^R, and active fractions are combined. The combined fractions are concentrated and the residue is digested in 10 ml of the upper phase of a solvent system consisting of n-heptane, benzene, acetone, isopropanol, 0.05 M, pH
35 7.0 aqueous phosphate buffer (5:10:3:2:5, v/v/v/v/v), and chromatographed on an Ito Coil Planet Centrifuge in the same system. Closely eluting active fractions are combined, concentrated and partitioned between methylene chloride and dilute ammonium hydroxide

(pH 9.0). The methylene chloride layer is separated and concentrated to yield the desired product as a white foam.

Example 4: Construction and characterization of *Streptomyces violaceoniger* NRRL 2834(pXC4) that produces 5-des-chalcosyl-5-desosaminoyl lankamycin

A. Construction of plasmid pKC4 and intermediates (see FIG. 12)

i) Construction of plasmid pKC1: The 2.4 kb DNA segment that carries the *eryCII* and *eryCIII* genes, comprised between coordinates 33 and 2413 of the sequence presented in SEQ ID NO:1, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:51 (5'-GAATGCATCTGGCTGGGCGGAGGGAATTCATG-3') and SEQ ID NO:52 (5'-GATCTAGACTTAAGTCATCGTGGTTCTCTCCTTCCTGC GGC-3'). After digestion with *NsiI* and *XbaI* the purified PCR fragment is ligated into *NsiI* and *XbaI* digested pK1 to give plasmid pKC1, 5.5 kb in size. The identity of plasmid pKC1, after transformation and isolation from *E. coli*, is verified by *EcoRI* digestion which releases fragments of 2.2 kb and 3.3 kb.

ii) Construction of plasmid pKC2: The 732 bp DNA segment that carries the *eryCVI* gene, comprised between coordinates 2331 and 3063 of the sequence presented in SEQ ID NO:2, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:53 (5'-GATCCTTAAGCTCCGGAGGGAGCAGGGATG-3') and SEQ ID NO:54 (5'-GATCTAGACCTAGGTCATCCGCGCACACCGACGAAC-3'). After digestion with *AflIII* and *XbaI* the purified PCR fragment is ligated into *AflIII* and *XbaI* digested pKC1 to give plasmid pKC2, 6.2 kb in size. The identity of plasmid pKC2, after transformation and isolation from *E. coli*, is verified by *XbaI-EcoRI* digestion which releases fragments of 0.95 kb, 2.2 kb and 3.1 kb.

iii) Construction of plasmid pKC3: The 2.7 kb DNA segment that carries the *eryCIV* and *eryCV* genes, comprised between coordinates 4650 and 7386 of the sequence presented in SEQ ID NO:2, is amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:55 (5'-GATCCTAGGCCGTCTACACCAGGACCGCCGG-3') and SEQ ID NO:56 (5'-GATCTAGATTAATCACCTTCCGCGCAGGAAGCCGC-3'). After digestion with *AvrII* and *XbaI* the purified PCR fragment is ligated into *AvrII* and *XbaI* digested pKC2 to yield plasmid pKC3, 9.0 kb in size. The identity of plasmid pKC3, after transformation and isolation from *E. coli*, is verified by *SphI* digestion which releases fragments of 4.0 kb and 5.0 kb.

iv) Construction of plasmid pKC4: The DNA sequence of the *eryCI* gene has been determined (GenBank Accession #X15541). The 1.1 kb DNA segment that carries the *eryCI* gene, comprised between coordinates 38 and 1161 of the sequence indicated above, is

amplified by PCR employing two deoxyoligonucleotides, SEQ ID NO:57 (5'-GATCTTAAG-CCGCCACTCGAACGGACACTCG-3') and SEQ ID NO:58 (5'-GATCTAGATCAAGCCC-CAGCCTTGAGGG-3'). After digestion with *MseI* and *XbaI* the fragment is ligated into *MseI* and *XbaI* digested pKC3 to give plasmid pKC4, 10.1 kb in size. The identity of plasmid pKC4, after transformation and isolation from *E. coli*, is verified by *KpnI* digestion which releases fragments of 0.15 kb, 0.31 kb, 4.1 kb and 5.5 kb. Plasmid pKC4 carries all of the *eryC* genes, *eryCI-eryCVI*, that are involved in the biosynthesis of desosamine and its attachment to the polydetide.

B. Construction of Plasmid pXSC4 (see FIG. 13): The 6.9 kb *NsiI-XbaI* segment of pKC4 that carries all of the *eryC* genes is isolated and ligated into *PstI-XbaI* digested pASX2, prepared as described in Example 2(A), to give the desired plasmid pXSC4, 14.9 kb in size, wherein all of the *eryC* genes are transcriptionally linked downstream of the *ermE** promoter on an *E. coli-Streptomyces* shuttle plasmid. The identity of plasmid pXSC4, after transformation and isolation from *E. coli*, is verified by the observation of fragments of 0.29 kb, 2.2 kb, and 12.4 kb after *EcoRI* digestion .

C. Construction of Plasmid pXC4 (see FIG. 13): The 6.9 kb *NsiI-XbaI* segment of pKC4 that carries all of the *eryC* genes is isolated and ligated into *NsiI-XbaI* digested pX1, prepared as described in Example 3(C)(ii), to give the desired plasmid pXC4, 11.5 kb in size, wherein all of the *eryC* genes are transcriptionally linked downstream of the *ermE** promoter on a *Streptomyces* plasmid. After transformation and isolation of the plasmid from *Streptomyces lividans* 1326, the identity of plasmid pXC4 is verified by the observation of fragments of 0.29 kb, 2.2 kb, and 9.0 kb after *EcoRI* digestion.

D. Construction of *Streptomyces violaceoniger* NRRL 2834(pXC4): Approximately 500 µg of the plasmid pXC4, isolated from *Streptomyces lividans* 1326(pXC4) , are electroporated into the lankamycin producer *Streptomyces violaceoniger* NRRL 2834 and several of the resulting Thio^R colonies that appear on the R3M-agar plates containing thiostrepton are analyzed for their plasmid content. The presence of plasmid pXC4 is verified by the observation of fragments of 0.29 kb, 2.2 kb, and 9.1 kb in size after *EcoRI* digestion of the plasmid.

E. Isolation, purification, and properties of 5-des-chalcosyl-5-desosaminoyl lankamycin: *S. violaceoniger* NRRL 2834(pXC4) is fermented for 5 days in SCM media with thiostrepton selection as described in General Methods. The fermentation broth is then cooled to 4°C and adjusted to pH 4.0 and extracted once with methylene chloride. The aqueous layer is readjusted to pH 9.0 and extracted twice with methylene chloride and the combined extracts are concentrated to a solid residue. This is digested in methanol and chromatographed over a column of Sephadex LH-20 in methanol. Fractions are tested for bioactivity against a sensitive organism, such as *Staphylococcus aureus* Th^R, and active

fractions are combined. The combined fractions are concentrated and the residue is digested in 10 ml of the upper phase of a solvent system consisting of n-heptane, benzene, acetone, isopropanol, 0.05 M, pH 7.0 aqueous phosphate buffer (5:10:3:2:5, v/v/v/v/v), and chromatographed on an Ito Coil Planet Centrifuge in the same system. Active fractions are combined, concentrated and partitioned between methylene chloride and dilute ammonium hydroxide (pH 9.0). The methylene chloride layer is separated and concentrated to yield the desired product as a white foam.

Although the present invention is illustrated in the examples listed above in terms of preferred embodiments, these examples are not to be regarded as limiting the scope of the invention. The above illustrations serve to describe the principles and methodologies involved in creating the types of genetic alterations that can be introduced into *Sac. erythraea* and/or other *Streptomyces* that result in the synthesis of novel glycosylation-modified polyketide products. Although a single Type I alteration, leading to the production of for example, 4"-deoxy-4"-oxo-erythromycin A, is specified herein, it is obvious to those skilled in the art that other Type I changes can be introduced into the *eryB* and/or *eryC* genes leading to novel glycosylation-modified polyketide structures. Examples of additional Type I alterations leading to useful novel compounds include but are not limited to: mutations in the *eryBVII* gene conceivably leading to 3- α -D-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B and mutations in the *eryCVI* gene conceivably leading to N-3 α' -des-dimethyl erythromycin A. Moreover, it is obvious that Type I alterations in two or more different *eryB* and/or *eryC* genes can be combined leading to novel glycosylation-modified polyketide structures. Examples of combinations of two Type I alterations leading to useful compounds include but are not limited to: mutations in the *eryBIV* and *eryBVII* genes conceivably leading to 3- α -D-4"-deoxy-4"-oxo-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B; mutations in the *eryBIV* and *eryCVI* genes conceivably leading to 4"-deoxy-4"-oxo-(N-3 α' -des-dimethyl)-erythromycin A; and mutations in the *eryBIV*, *eryBVII*, and *eryCVI* genes conceivably leading to 3- α -D-4"-deoxy-4"-oxo-mycarosyl-5- β -D-(N-3 α' -des-dimethyl)-desosaminoyl-12-hydroxy-erythronolide B. All Type I mutations or combinations of two or more Type I mutations in the *eryBII*, *eryBIV*, *eryBV*, *eryBVI*, *eryBVII*, *eryCII*, *eryCIII*, *eryCIV*, *eryCV*, or *eryCVI* genes, the *Sac. erythraea* strains that carry said mutations or combinations of mutations, and the corresponding polyketides produced from said strains, therefore, are included within the scope of the present invention.

Although the Type II mutation specified herein was constructed with the *eryBVII* gene on a self-replicating plasmid it is obvious that other *eryB* genes and *eryC* genes can be expressed in an antisense orientation leading to novel glycosylation-modified polyketide structures. Examples of additional Type II alterations leading to useful compounds include but are not limited to: antisense expression of the *eryBIV* gene conceivably leading to 4"-

deoxy-4"-oxo-erythromycin A and antisense expression of the *eryCVI* gene conceivably leading to N-3 α '-des-dimethyl erythromycin A. Moreover, it will occur to those skilled in the art that promoters other than the *ermE** promoter, for example the *melC* promoter of pIJ702, will be suitable for antisense expression, and that many self-replicating vectors in addition to pWHM4 will function to carry the antisense alteration. It will also occur to those skilled in the art that a self-replicating vector is not required for this invention and that the antisense alteration can be introduced directly into the chromosome using the same principles employed to construct a Type I gene alteration. An example of a Type II alteration that is introduced directly into the chromosome is the *eryBVII* antisense alteration described in Example 2 wherein DNA segments immediately upstream of the *eryK* gene are used to flank the *ermE-eryBVII*-phage fd terminator grouping in a pWHM3 vector, and this vector is integrated into and then resolved from the chromosome leaving the *ermE*-eryBVII*-phage fd terminator grouping stably incorporated into this nonessential region of the chromosome of *Sac. erythraea* conceivably leading to the production of 3- α -D-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B. All Type II mutations in the *eryBII*, *eryBIV*, *eryBV*, *eryBVI*, *eryBVII*, *eryCII*, *eryCIII*, *eryCIV*, *eryCV*, or *eryCVI* genes whether carried on a self-replicating plasmid or integrated into a nonessential region of the chromosome, the *Sac. erythraea* strains that carry said mutations, and the corresponding polyketides produced from said strains, therefore, are included within the scope of the present invention.

Although Type III alterations, leading to the production of 5-des-chalcosyl-5-desosaminoyl lankamycin in *Streptomyces violaceoniger* and 3-des-oleandrosyl-3-mycarosyl oleandomycin in *Streptomyces antibioticus*, are specified herein, it is obvious that Type III alterations can be introduced into any polyketide producing microorganism leading to novel glycosylation modified polyketides. It will also occur to those skilled in the art that both the *eryB* and *eryC* genes can either be cotransformed into a polyketide producing microorganism or grouped together on a single vector that is introduced into a polyketide producing microorganism. An example of a Type III change using both the *eryB* and *eryC* genes together is their introduction into *Streptomyces violaceoniger* conceivably leading to 3-des-(4"-O-acetylarcanosyl)-3-mycarosyl-5-des-chalcosyl-5-desosaminoyl lankamycin. Although the Type III alterations specified herein have indicated a specific genetic order of the *eryB* or *eryC* genes, it will occur to those skilled at the art that many different genetic arrangements of the *eryB* or *eryC* genes will produce similar results. It will also that occur to those skilled at the art that certain arrangements of the *eryB* and/or *eryC* genes that lack one or more of the respective *eryB* and/or *eryC* genes will lead to the production of novel glycosylated polyketides in which intermediate compounds in the biosynthesis of mycarose and/or desosamine, respectively, such as those outlined in FIGS. 2 and 3, are attached to the polyketide. An example of a Type III alteration in which only a subset of the *eryB* and/or

eryC genes are used is the introduction of a pXC4 derivative that lacks the *eryCVI* gene, removed by digestion of plasmid pXC4 with *Afl*III and *Avr*II followed by treatment with the Klenow fragment of DNA polymerase I and religation, into *Streptomyces violaceoniger* leading to the production of to 5-des-chalcosyl-5-(N-3 α '-des-dimethyl desosaminoyl) lankamycin. It will also that occur to those skilled at the art that promoters other than *ermE* or *ermE**, such as the *melC* promoter of plasmid pIJ702, and vectors other than pWHM4 or pIJ702 can also be utilized in the construction of a Type III alteration, and these variants are, of course, considered to be within the scope of the invention. Finally, it will also occur to those skilled in the art that a self-replicating vector is not required for this invention and that an assembly of sugar biosynthesis genes can be introduced directly into the chromosome of a heterologous host using the same principles employed to construct a Type I gene alteration once a nonessential region of the heterologous host chromosome has been identified. Alternatively, plasmids or bacteriophages which undergo site-specific recombination with host genes may also be used to introduce *eryB* and *eryC* genes into a host to effect Type III alterations. All Type III alterations using one or more of the *eryBII*, *eryBIV*, *eryBV*, *eryBVI*, *eryBVII*, *eryCII*, *eryCIII*, *eryCIV*, *eryCV*, or *eryCVI* genes, the polyketide producing strains that carry said alterations, and the corresponding polyketides produced from said strains, therefore, are included within the scope of the present invention.

In addition, it is also possible to create combinations of Type I and Type II alterations such that some Type I *eryB* and/or *eryC* mutations are introduced directly into the *Sac. erythraea* chromosome in the appropriate locus, while other *eryB* and/or *eryC* genes are inactivated by Type II alterations using a self-replicating or integrating vector. For example, combination of a Type I alteration, such as a mutation in *eryBIV*, and a Type II alteration, such as transformation with pASBVII, will conceivably lead to production of 3- α -D-4"-deoxy-4"-oxo-mycarosyl-5- β -D-desosaminoyl-12-hydroxy-erythronolide B. All combinations of two or more alterations of Type I and Type II, the *Sac. erythraea* strains that carry such alterations, and the glycosylated polyketides produced from such strains are included within the scope of the present invention.

As an extension of the examples reported with the *eryB* and/or *eryC* genes, it is possible to apply the method described herein to heterologous sugar biosynthesis genes that are similar to the *eryB* and/or *eryC* genes. The construction of strains carrying heterologous sugar biosynthesis genes that lead to the production of novel glycosylated polyketides requires: (i) cloning of the sugar biosynthesis genes from any other glycosylated-polyketide producing actinomycete, (ii) determining the nucleotide sequence of the cloned gene(s); (iii) excising and assembling the cloned gene(s) into vectors suitable for Type I, Type II, or Type III alterations; and (iv) transformation of polyketide producing microorganisms and screening for the novel compound. Any polyketide-associated sugar biosynthesis gene can thus be

precisely excised from the genome of a glycosylated polyketide producing microorganism and altered or arranged with other sugar biosynthesis genes and then introduced into the same or another polyketide producing microorganism to create a novel glycosylated polyketide of predicted structure. Thus, for example, a Type I or Type II alteration of a heterologous gene that is similar to an *eryB* and/or *eryC* gene, such as can be found in the *eryBVII* homolog for the synthesis of L-oleandrose in *Streptomyces antibioticus*, to result in the production of 3-des-L-oleandrosyl-3-D-oleandrosyl oleandomycin is included within the scope of the present invention. Similarly, a Type III assembly of the genes for the synthesis of a sugar other than mycarose or desosamine, such as can be found in the genes for the synthesis of angolosamine in *Streptomyces eurythermus*, and their transformation into *Sac. erythraea* to result in the synthesis of 5-des-desosaminoyl-5-angolosaminoyl-erythromycin A is included within the scope of the present invention.

It will occur to those skilled in the art that the Type I, Type II, and Type III genetic manipulations described herein and the polyketide producing microorganisms into which they are introduced are in no way exclusive. Hence, the choice of a convenient host and the choice of a Type I, Type II, or Type III alteration is based solely on the relatedness of the desired novel glycosylated polyketide to a natural counterpart. Therefore, Type I, Type II, and Type III alterations can be constructed in any polyketide producing microorganism employing either endogenous or exogenous sugar biosynthesis genes. Thus all Type I, Type II, and Type III mutations or various combinations thereof constructed in any polyketide producing microorganism according to the principles described herein, and the respective polyketides produced from such strains, are included within the scope of the present invention. Examples of glycosylated polyketides that can be altered by creating Type I, Type II, or Type III changes in the producing microorganisms include, but are not limited to macrolide antibiotics such as erythromycin, tylosin, spiramycin, etc; aromatic polyketides such as daunorubicin and doxorubicin, etc; polyenes such as candicidin, amphotericins, etc; and other complex polyketides such as avermectin.

Whereas the novel derivatives or modifications of erythromycin described herein have been specified as the A derivatives, such as 4"-deoxy-4"-oxo-erythromycin A, those skilled in the art understand that the wild type strain of *Sac. erythraea* produces a family of erythromycin compounds, including erythromycin A, erythromycin B, erythromycin C, and erythromycin D. Thus, modified strains of *Sac. erythraea*, such as strain ERBIV, for example, would be expected to produce the corresponding members of the 4"-deoxy-4"-oxo-erythromycin family, including 4"-deoxy-4"-oxo-erythromycin A, 4"-deoxy-4"-oxo-erythromycin B, 4"-deoxy-4"-oxo-erythromycin C, and 4"-deoxy-4"-oxo-erythromycin D. Similarly, all other modified strains of *Sac. erythraea* that produce novel glycosylated erythromycin derivatives would be expected to produce the A, B, C, and D forms of said

derivatives. For example, modified *Sac. erythraea* strains that produce 6-deoxyerythromycin, 6,12-dideoxyerythromycin and 6,7-anhydroerythromycin would be expected to produce novel glycosylation-modified polyketides by introduction of the additional modification of a Type I, II or III change in a sugar biosynthesis gene. Therefore, all members of the family of each of the novel erythromycins described herein or produced by these methods are included within the scope of the present invention.

Variations and modifications of the methods for obtaining the desired plasmids, hosts for cloning and choices of vectors and *eryB* and/or *eryC* genes to clone and modify, other than those described herein will occur to those skilled in the art. For example, although we have described the use of plasmids pWHM3, pWHM4, and pIJ702, other vectors can be employed wherein all or part of said plasmids is replaced by other DNA segments that function in a similar manner, such as replacing the pUC19 component of pWHM3 and pWHM4 with pBR322, available from BRL; or employing different segments of the pIJ101 replicon in pWHM3 and pIJ702, or the pJV1 replicon in pWHM4, respectively; or employing selectable markers other than thiostrepton- or ampicillin-resistance. These are just a few of a long list of possible examples all of which are included within the scope of the present invention. Similarly, the segments of the *eryB* and *eryC* loci that have been specified herein to generate the various Type I, Type II, and Type III alterations can readily be substituted for other segments of different length encoding the same functions, either produced by PCR-amplification of genomic DNA or of an isolated clone, or by isolating suitable restriction fragments from *Sac. erythraea*. In the same way it is possible to create Type I mutations functionally equivalent to those described herein by altering through deletion, insertion, or site directed mutagenesis different portions of the corresponding genes. It is also possible to create Type II mutations functionally equivalent to those described herein by employing larger or smaller portions of the corresponding genes; and it is possible to create Type III mutations using larger or smaller segments of the corresponding genes in the same or different linear order described herein. Additional modifications include changes in the restriction sites used for cloning or in the general methodologies described above. All such changes are included in the scope of the present invention. It will also occur to those skilled in the art that different methods are available to ferment *Sac. erythraea* and other polyketide producing microorganisms and to extract the novel polyketides specified herein, and all such methods are also included within the scope of this invention.

It will also be apparent that many modifications and variations of the invention as set forth herein are possible without departing from the spirit and scope thereof, and that, accordingly, such limitations are imposed only as indicated by the appended claims.

We claim:

1. An isolated single or double stranded polynucleotide having a nucleotide sequence which comprises (a) a nucleotide sequence selected from the group consisting of (i) the sense sequence of SEQ ID NO:1 from about nucleotide position 54 to about nucleotide position 1136; (ii) the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412; (iii) sense sequence of SEQ ID NO:1 from about nucleotide position 2409 to about nucleotide position 3410; (iv) the sense sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048; (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295; (vi) the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061; (vii) the sense sequence of SEQ ID NO:2 from about nucleotide position 3214 to about nucleotide position 4677; (viii) the sense sequence of SEQ ID NO:2 from about nucleotide position 4674 to about nucleotide position 5879; (iv) the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386; and (x) the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996;
 - (b) sequences complementary to the sequences of (a);
 - (c) sequences that, on expression, encode a polypeptide encoded by the sequences of (a); and
 - (d) analogous sequences that hybridize under stringent conditions to the sequences of (a).
2. The polynucleotide of claim 1 that is a DNA molecule or RNA molecule.
3. The polynucleotide of claim 2 wherein the nucleotide sequence is the nucleotide sequence of (a) selected from the group consisting of (i) the sense sequence of SEQ ID NO:1 from about nucleotide position 54 to about nucleotide position 1136; (ii) the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412; (iii) the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061; (iv) the sense sequence of SEQ ID NO:2 from about nucleotide position 4674 to about nucleotide position 5879; and (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386.
4. The polynucleotide of claim 2 wherein the nucleotide sequence is the nucleotide sequence of (a) selected from the group consisting of (i) sense sequence of SEQ ID NO:1 from about nucleotide position 2409 to about nucleotide position 3410; (ii) the sense

- sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048; (iii) the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295; (iv) the sense sequence of SEQ ID NO:2 from about nucleotide position 3214 to about nucleotide position 4677; and (v) the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996.

5. The polynucleotide of claim 2 wherein the nucleotide sequence is the nucleotide sequence of (a) having the sense sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048.
6. A vector comprising the DNA molecule of claim 2.
7. The vector of claim 6 further comprising an enhancer-promoter operatively linked to the polynucleotide.
8. The vector of claim 6 wherein the polynucleotide has the nucleotide sequence of claim 5.
9. A host cell transformed with the vector of claim 6 or claim 7 or claim 8.
10. The transformed host cell of claim 9 that is a bacterial cell.
11. The transformed host cell of claim 10 wherein the bacterial cell is selected from the group consisting of *Streptomyces* and *E. coli*.
12. A method for directing the biosynthesis of specific glycosylation-modified polyketides by genetic manipulation of a polyketide-producing microorganism, said method comprising the steps of:
 - (1) isolating a sugar biosynthesis gene-containing DNA sequence according to claim 1;
 - (2) identifying within said gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to a polyketide;
 - (3) creating one or more specified changes into said DNA fragment or fragments, thereby resulting in an altered DNA sequence;
 - (4) introducing said altered DNA sequence into a polyketide-producing microorganism to replace the original sequence, said altered DNA sequence, when translated,

resulting in altered enzymatic activity capable of effecting the production of said specific glycosylation-modified polyketide;

- 15 (5) growing a culture of said altered polyketide-producing microorganism under conditions suitable for the formation of said specific glycosylation-modified polyketide; and
(6) isolating said specific glycosylation-modified polyketide from said culture.

13. The method of claim 12 wherein said specified change in said DNA fragment or fragments results in the inactivation of at least one enzymatic activity involved in the biosynthesis of a polyketide-associated sugar or in its attachment to a polyketide.

14. The method of claim 13 wherein said polyketide-associated sugar is L-mycarose.

15. The method of claim 13 wherein said polyketide-associated sugar is D-desosamine.

16. A method for directing the biosynthesis of specific glycosylation-modified polyketides by genetic manipulation of a polyketide-producing microorganism, said method comprising the steps of:

- 5 (1) isolating a sugar biosynthesis gene-containing DNA sequence according to claim 1;

(2) identifying within said gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to a polyketide;

- 10 (3) reversing the strand orientation of said DNA fragment or fragments, thereby resulting in an altered DNA sequence which, when transcribed, results in production of an antisense mRNA;

(4) introducing said altered DNA sequence into a polyketide-producing microorganism having an mRNA capable of binding to said antisense mRNA to produce an altered polyketide-producing microorganism capable of producing said specific
15 glycosylation-modified polyketide;

- (5) growing a culture of said altered polyketide-producing microorganism under conditions suitable for the formation of said specific glycosylation-modified polyketide; and
(6) isolating said specific glycosylation-modified polyketide from said culture.

17. A method for directing the biosynthesis of specific glycosylation-modified polyketides by genetic manipulation of a polyketide-producing microorganism, said method comprising the steps of:

- (1) isolating a sugar biosynthesis gene-containing DNA sequence according to claim

5 1;

(2) identifying within said gene-containing DNA sequence one or more DNA fragments responsible for the biosynthesis of a polyketide-associated sugar or its attachment to a polyketide;

10 (3) introducing said DNA fragment or fragments into a distinct polyketide-producing microorganism to produce an altered polyketide-producing microorganism capable of producing said specific glycosylation-modified polyketide;

(4) growing a culture of said polyketide-producing microorganism containing said DNA fragment or fragments under conditions suitable for the formation of said specific glycosylation-modified polyketide; and

15 (6) isolating said specific glycosylation-modified polyketide from said culture.

18. The method of claim 13 or claim 16 or claim 17 wherein said DNA fragment comprises one or more genes which encode an enzymatic activity involved in the biosynthesis of L-mycarose or in its attachment to a polyketide.

19. The method of claim 13 or claim 16 or claim 17 wherein said DNA fragment comprises one or more genes which encode an enzymatic activity involved in the biosynthesis of D-desosamine or in its attachment to a polyketide.

20. The method of claim 13 or claim 16 or claim 17 wherein said DNA fragment is the sequence of claim 8.

21. An isolated polypeptide having an amino acid sequence encoded by a nucleotide sequence selected from the group consisting of the sense sequence of SEQ ID NO:1 from about nucleotide position 54 to about nucleotide position 1136; the sense sequence of SEQ ID NO:1 from about nucleotide position 1147 to about nucleotide position 2412; sense sequence
5 of SEQ ID NO:1 from about nucleotide position 2409 to about nucleotide position 3410; the sense sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048; the sense sequence of SEQ ID NO:2 from about nucleotide position 1048 to about nucleotide position 2295; the sense sequence of SEQ ID NO:2 from about nucleotide position 2348 to about nucleotide position 3061; the sense sequence of SEQ ID NO:2 from
10 about nucleotide position 3214 to about nucleotide position 4677 ; the sense sequence of SEQ ID NO:2 from about nucleotide position 4674 to about nucleotide position 5879; the sense sequence of SEQ ID NO:2 from about nucleotide position 5917 to about nucleotide position 7386; and the sense sequence of SEQ ID NO:2 from about nucleotide position 7415 to about nucleotide position 7996.

22. An isolated polypeptide of claim 31 encoded by the sequence of SEQ ID NO:2 from about nucleotide position 80 to about nucleotide position 1048.

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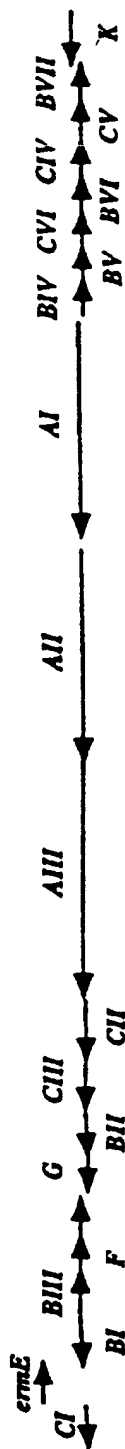


FIG. 1A

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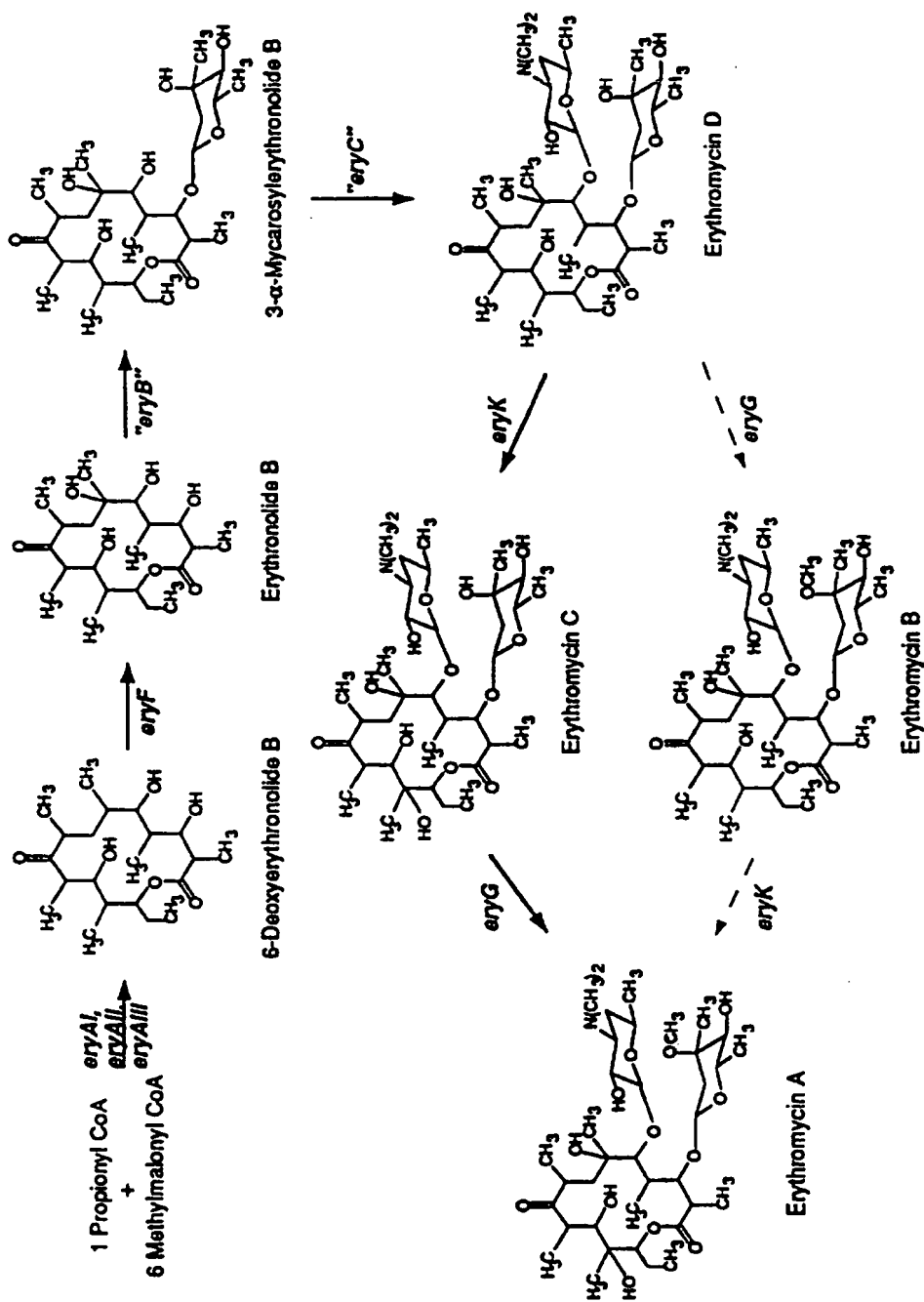


FIG. 1B

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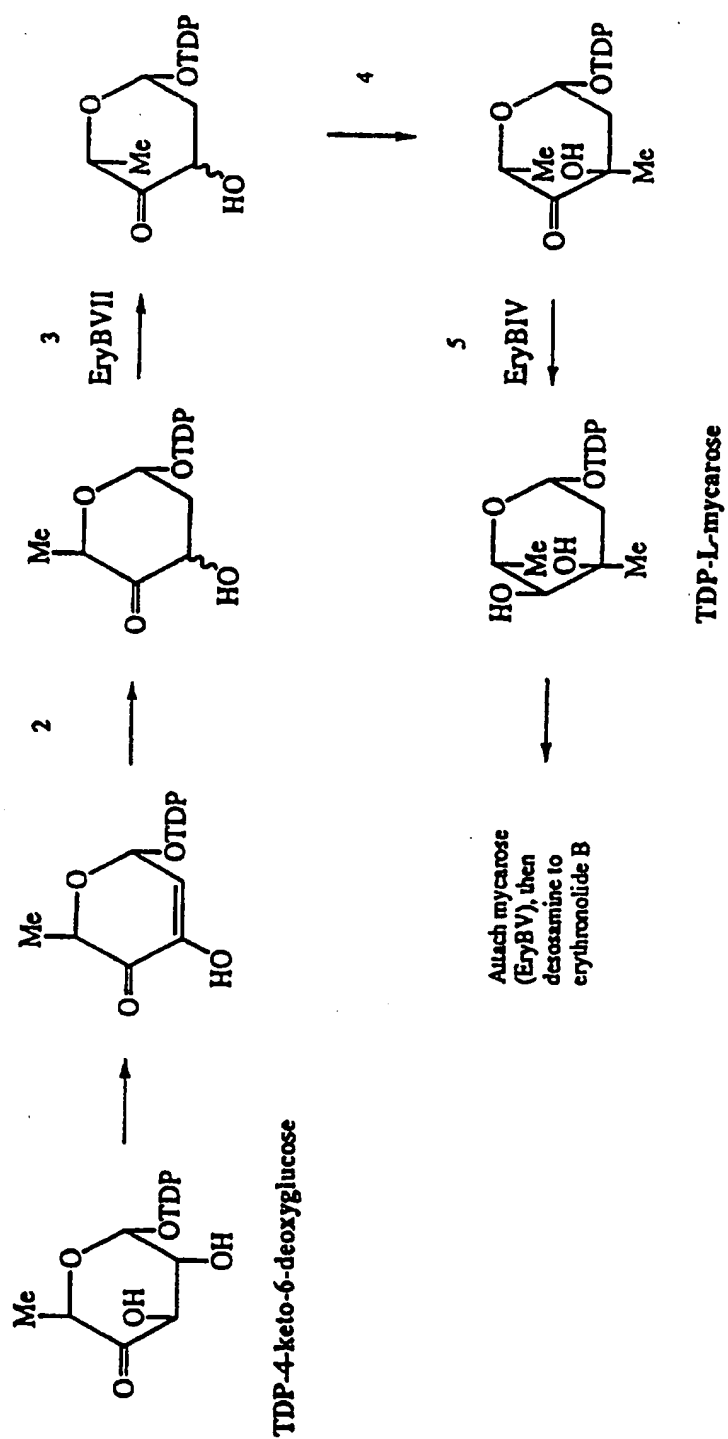


FIG. 2

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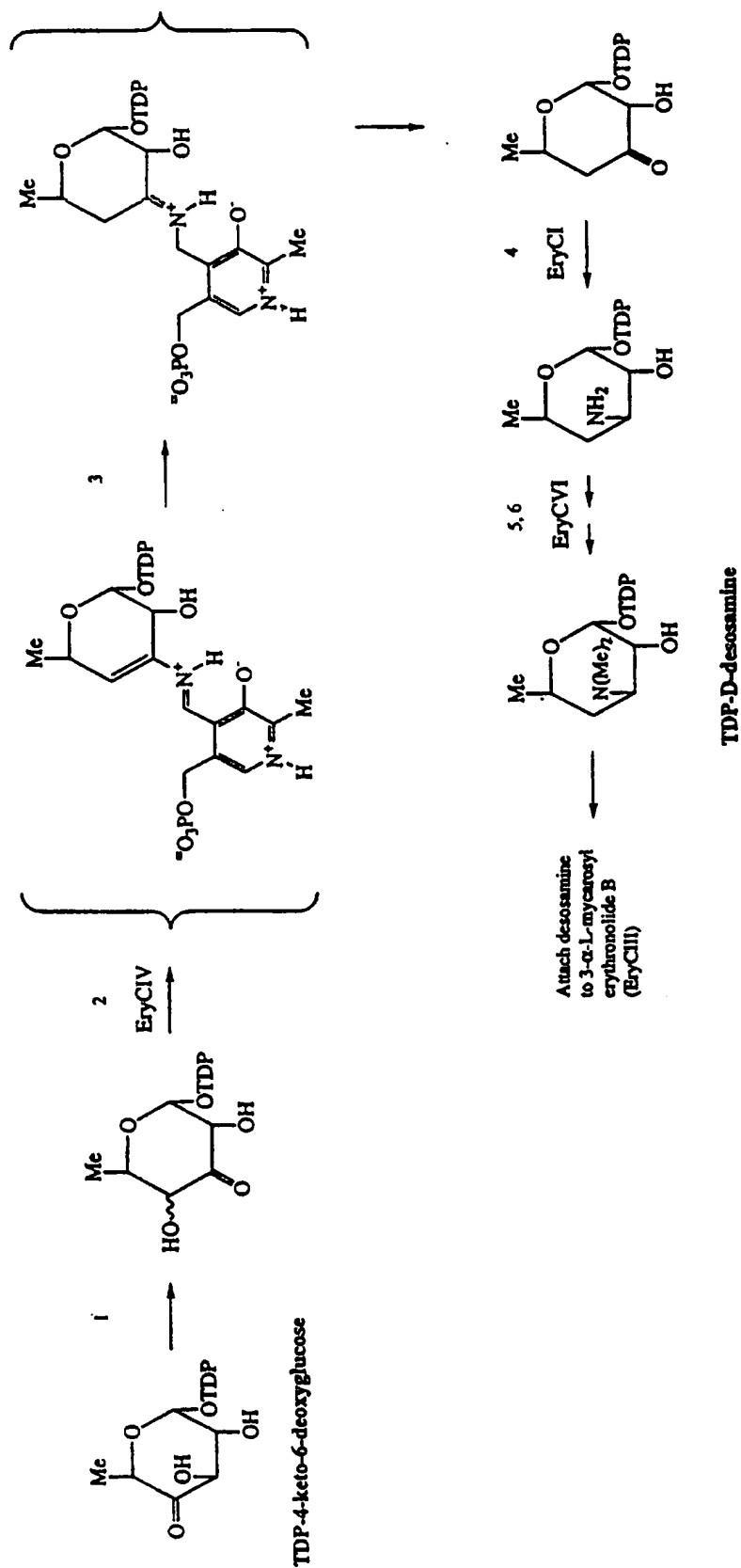


FIG. 3

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1 CACGCCGACGCGATCGCGCGGCACATCGACGCCTGGCTGGGCGGAGGGAATTCATGACCA 60
M T T

61 CGACCGATCGCGCCGGGCTGGGCAGGCAGCTCCAGATGATCCGCGGCCTGCACTGGGGTT 120
T D R A G L G R Q L Q M I R G L H W G Y

121 ACGGCAGCAACGGCGACCCTTACCCGATGCTGCTGTGCGGACACGACGACGACCCGCAGC 180
G S N G D P Y P M L L C G H D D D P Q R

181 GCCGGTACCGCTCGATGCGCGAGTCCGGTGTGCGGCGCAGGACCGAGACGTGGGTGGTGG 240
R Y R S M R E S G V R R R T E T W V V A

241 CCGACCACGCCACCGCCCGGCAGGTGCTCGACGACCCCGCGTTACCCGCGCCACCGGAC 300
D H A T A R Q V L D D P A F T R A T G R

301 GCACACCGGAATGGATGCGGGCCGCGGGCGGCCACCCGCGGAGTGGGCCCAGCCGTTCC 360
T P E W M R A A G A P P A E W A Q P F R

361 GGGACGTGCACGCCCGTCTTGGGAAGCGAGGTCCCCGACGTGCGGGAACCTGGCGGAGA 420
D V H A A S W E G E V P D V G E L A E S

421 GCTTCGCCGCTCTGCTCCCCGGCGCGGGCGCGCGCTGGACCTGGTGGGCGACTTCGCCT 480
F A G L L P G A G A R L D L V G D F A W

481 GGCAGGTACCGGTGCAGGGCATGACCGCCGTGCTCGGCGCAGCCGAGTGTCTGCGCGGCG 540
Q V P V Q G M T A V L G A A G V L R G A

541 CCGCGTGGGACGCCCGCGTCCAGCCTGGACGCCAGCTCAGCCCGCAGCAGCTCGCGGTGA 600
A W D A R V S L D A Q L S P Q Q L A V T

601 CCGAAGCAGCGGTGCGGGCACTGCCCGCCGACCCGCACTGCGCGCCCTGTTCGCCGGGG 660
E A A V A A L P A D P A L R A L F A G A

661 CCGAGATGACCGCGAACACCGTGGTGCAGCGCGTCTGGCCGTCTCGGCCGAACCGGGG 720
E M T A N T V V D A V L A V S A E P G L

721 TGGCCGAACGGATCGCCGACGACCCCGCCCGCGCAGCGAACCGTCCGCCGAGGTGCTGC 780
A E R I A D D P A A A Q R T V A E V L R

781 GCCTGCACCCGGCATTGCACCTGGAGCGGCGCACGGCCACCGCAGAGGTGCGGCTCGGCG 840
L H P A L H L E R R T A T A E V R L G E

841 AGCACGTGATCGGCGAAGGCGAGGAGTGTGTCGTCGTCGCGGCGGCCAACCGCGACC 900
H V I G E G E E V V V V V A A A N R D P

901 CGGAGGTCTTCGCCGAGCCCGACCGCCTGACGTGGACCGCCCCGACCGCGACCGCGCGC 960
E V F A E P D R L D V D R P D A D R A L

FIG. 4A-1

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961 TGTCGGCACATCGCGGCCACCCGGCAGGCTGGAGGAGCTGGTCACCGCGCTCGCCACCG 1020
S A H R G H P G R L E E L V T A L A T A

1021 CCGCACTGCGGGCCGCGGCCAAGGCGCTGCCCGGACTCAAGCCAGCGGCCCGGTCGTCC 1080
A L R A A A K A L P G L T P S G P V V R

1081 GCGCGCCCGGATCACCCGTCTTGGGGGAACCAACCGCTGCCCCGTGAGCTCTGAGGAT 1140
R R R S P V L R G T N R C P V E L *

1141 TCCGCGATGCGCGTCGTCTTCTCCTCCATGGCCAGCAAGAGCCACCTCTTCGGCCTCGTC 1200
M R V V F S S M A S K S H L F G L V

1201 CCCCTCGCATGGGCGTTCCGCGCGCGGGGCACGAGGTCCGCGTGGTCGCGTCCCCGGCG 1260
P L A W A F R A A G H E V R V V A S P A

1261 CTCACCGAGGACATCACCGCGGGCGGCTGACCGCCGTCCCGGTGGCAACCGACGTCGAC 1320
L T E D I T A A G L T A V P V G T D V D

1321 CTCGTGGACTTCATGACCCACGCGGGCCACGACATCATCGACTACGTCCGGAGCCTGGAC 1380
L V D F M T H A G H D I I D Y V R S L D

1381 TTCAGCGAGCGGGACCCCGCCACCTTGACCTGGGAGCACCTGCGGGGCATGCAGACCGTG 1440
F S E R D P A T L T W E H L R G M Q T V

1441 CTCACCCCGACCTTCTACGCCCTGATGAGCCCGGACACGCTCATCGAAGGCATGGTCTCG 1500
L T P T F Y A L M S P D T L I E G M V S

1501 TTCTGCCGGAAGTGGCGGCCCGACCTGGTCATCTGGGAGCCGCTCACCTTCGCCGCGCCC 1560
F C R K W R P D L V I W E P L T F A A P

1561 ATCGCGGGCGCGGTGACCGGAACGCGCACGCGCGGCTGCTGTGGGGACCCGACATCAC 1620
I A G A V T G T P H A R L L W G P D I T

1621 ACCCGGGCGCGGCAGAACTTCTCGGCCTGCTGCCCCGACCAGCCGGAGGAGCACCGGGAG 1680
T R A R Q N F L G L L P D Q P E E H R E

1681 GGCCCGCTCGCCGAGTGGCTCACCTGGACGCTGGAGAAGTACGGCGGCCCGGCCTTCGAC 1740
G P L A E W L T W T L E K Y G G P A F D

1741 GAGGAGGTGGTCGTGCGGCAGTGGACGATCGACCCCGCCCGCGCGATCAGGCTCGAC 1800
E E V V V G Q W T I D P A P A A I R L D

1801 ACCGGCCTGAAGACCGTCGGGATGCGCTACGTCGACTACAACGGGCGCTCCGTGGTGCCG 1860
T G L K T V G M R Y V D Y N G P S V V P

1861 GAATGGCTGCACGACGAGCCGAGCGCCGCGCGTGTGCCCTCACGCTCGGGATCTCCAGC 1920
E W L H D E P E R R R V C L T L G I S S

FIG. 4A-2

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1921	CGCGAGAACAGCATCGGGCAGGTCTCCATCGAGGAGCTGCTGGGTGCCGTCGGCGACGTC	1980
	R E N S I G Q V S I E E L L G A V G D V	
1981	GACGCCGAGATCATCGCGACCTTCGACGCGCAGCAGCTAGAAGGCGTCGCGAACATCCCG	2040
	D A E I I A T F D A Q Q L E G V A N I P	
2041	CACAACGTCCGCACGGTCGGCTTCGTCCCGATGCACGCGCTGCTGCCGACCTGCGCGGCG	2100
	H N V R T V G F V P M H A L L P T C A A	
2101	ACGGTGCACACGCGGACCCGGGAGCTGGCACACCGCGGCGATCCACGGCGTGCCGCGAG	2160
	T V H H G G P G S W H T A A I H G V P Q	
2161	GTGATCCTGCCCCGACGGCTGGGACACCGGCGTGCGCGCGCAGCGCACGGAATTCCGGG	2220
	V I L P D G W D T G V R A Q R T Q E F G	
2221	GCGGGGATCGCGCTGCCCCGTGCCCCGAGCTGACCCCCGACCAGCTCCGGGAGTCGGTGAAG	2280
	A G I A L P V P E L T P D Q L R E S V K	
2281	CGGGTCCTCGACGACCCGGCCACCGCGCCGGCGCGGCGGATGCGCGACGACATGCTC	2340
	R V L D D P A H R A G A A R M R D D M L	
2341	GCGGAGCCGTCACCGGCCGAGGTCTGCGGCATCTGCGAGGAAGTGGCCGCGAGGAAGGAGA	2400
	A E P S P A E V V G I C E E L A A G R R	
2401	GAACCACGATGACCACCGACGCCGCGACGACAGTGGGCTCGGGCGTTCCGCGCTGCTCA	2460
	E P R * M T T D A A T H V R L G R S A L L T	
2461	CCAGCAGGCTCTGGCTCGGCACGGTGAACCTCAGCGGACGCGTCGAGGACGACGACGCGC	2520
	S R L W L G T V N F S G R V E D D D A L	
2521	TGCGCCTGATGGACCACGCCCGGGACCGGCGCATCAACTGCCTCGACACCGCCGACATGT	2580
	R L M D H A R D R G I N C L D T A D M Y	
2581	ACGGCTGGCGGCTCTACAAGGGCCACACCGAGGAGCTGGTGGGCAGGTGGCTGGCCCAGG	2640
	G W R L Y K G H T E E L V G R W L A Q G	
2641	GCGGCGGACGGCGCGAGGACACCGTGCTGGCGACCAAGGTGCGGCGGAGATGAGCGAGC	2700
	G G R R E D T V L A T K V G G E M S E R	
2701	GCGTCAACGACAGCGGGCTGTCGGCGCGGCACATCATCGCCTCCTGCGAGGGATCGCTGC	2760
	V N D S G L S A R H I I A S C E G S L R	
2761	GCAGGCTGGGCGTCGACCACATCGACGTCTACCAGATGCACCACATCGACCGGTCCGCGC	2820
	R L G V D H I D V Y Q M H H I D R S A P	
2821	CGTGGGACGAGGTGTGGCAGGCCATGGACAGCCTCGTCGCCAGCGGCAAGGTCTCCTACG	2880
	W D E V W Q A M D S L V A S G K V S Y V	

FIG. 4A-3

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2881 TCGGCTCGT[˙]CGAACTTCGCGGGCTGGCACATCGCCGCCG[˙]CGCAGGAGAACGCGCCCGCC[˙] 2940
G S S N F A G W H I A A A Q E N A A R R

2941 GCCACTCCCTGGGCATGGTCTCCACCACTGCCTGTACAACCTGGCGGT[˙]CCGGCAGCCG[˙] 3000
H S L G M V S H Q C L Y N L A V R H A E

3001 AGCTGGAGGT[˙]GCTGCCCCGCCGCGCAGGCCTACGGGCTCGGCGTCTTCGCCTGGTCGCCG[˙]C 3060
L E V L P A A Q A Y G L G V F A W S P L

3061 TGCACGGCGGCTGCTCAGCGGAGCGCTGGAGAAGCTGGCCGCGGGCACCGCGGTGAAGT[˙] 3120
H G G L L S G A L E K L A A G T A V K S

3121 CGGCGCAGGGCCGTGCGCAGGTGCTGTG[˙]CCGTCCCTGCGCCCGCGGAT[˙]CGAGGCCTACG[˙] 3180
A Q G R A Q V L L P S L R P A I E A Y E

3181 AGAAGTTCTGCCGCAACCTCGGCGAAGACCCGGCCGAGGTGGGGCTCGCATGGGTGCTGT[˙] 3240
K F C R N L G E D P A E V G L A W V L S

3241 CCGGCCCCGGCATCGCCGGCGCGTCA[˙]TGCGCCCGGAACCCCCGAGCAGCTCGACTCCG[˙] 3300
R P G I A G A V I G P R T P E Q L D S A

3301 CGCTGAAGGCGTCCGCGATGACCCTGGACGAGCAGGCGCTGTCCGAAC[˙]TGGACGAGATCT[˙] 3360
L K A S A M T L D E Q A L S E L D E I F

3361 TCCCCGCGGTGGCCTCCGGCGGGCGGGCGCGGAAGCCTGGTTGCAGTGAGCACAAGAGG[˙] 3420
P A V A S G G A A P E A W L Q *

3421 AACCGAGAAAGGATACGGCTGGTGAGCGTGAAGCAGAAGTCAGCGTTGCAGGACCTGGTC[˙] 3480

3481 GACTTCGCAAGTGGCACGTGTGGACCAGGGTGGCGCCGTCCAGCCGTGCGCGCTGGCC[˙] 3540

3541 TACGAGCTGTTCGCCGACGACCACGAGGCCACGACCGAGGGCGCCTACATCAACCTCGGC[˙] 3600

3601 TACTGGAAGCCCGGTGCGCGGCTGGAGGAGGCCAAC[˙]CAGGAGCTGGCGAACCAGCTC[˙] 3660

3661 GCCGAGGCCCGGGGATCAGCGAGGGCGACGAGGTGCTCGACGTCGGGTTCGGGCTCGGC[˙] 3720

3721 GCGCAGGACTTCTTCTGGCTCGACCTGCAGCCAGCT 3756

FIG. 4A-4

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1 CGGGTTGCCGCACATCGCGCTGGGGAGATTCTTTGAATTTGCCCCGTAGCACCGACCTGG 60

61 AAAGCGAGCAAATGCTCCGGTGAATGGGATCAGTGATTCCCGCGTCAATTGATCACCT 120
V N G I S D S P R Q L I T L

121 TCTGGGCGCTTCCGGCTTCGTGGGAGCGCGGTTCTGCGCGAGCTGCGCGACCAACCGGT 180
L G A S G F V G S A V L R E L R D H P V

181 CCGGCTGCGCGCGGTGTCCCGCGGCGGAGCGCCCGCGGTTCGCCCCGGCGCGGAGGT 240
R L R A V S R G G A P A V P P G A A E V

241 CGAGGACCTGCGCGCCGACCTGCTGGAACCGGGCCGGCCGCGCCGCGATCGAGGACGC 300
E D L R A D L L E P G R A A A A I E D A

301 CGACGTGATCGTGACCTGGTGGCGCACGCAGCGGGCGGTTCCACCTGGCGCAGCGCCAC 360
D V I V H L V A H A A G G S T W R S A T

361 CTCCGACCCGGAAGCCGAGCGGGTCAACGTGGGCTGATGCACGACCTCGTCCGCGCGCT 420
S D P E A E R V N V G L M H D L V G A L

421 GCACGATCGCCGAGGTGCGACGCGCCCGTGTGCTCTACGCGAGCACCGCACAGGCCGC 480
H D R R R S T P P V L L Y A S T A Q A A

481 GAACCCGTGCGCGGCCAGCAGGTACGCGCAGCAGAAGACCAGGGCCGAGCGCATCCTGCG 540
N P S A A S R Y A Q Q K T E A E R I L R

541 CAAAGCCACCGACGAGGGCCGGGTGCGCGCGTGATCCTGCGGCTGCCCCGCGTCTACGG 600
K A T D E G R V R G V I L R L P A V Y G

601 CCAGAGCGGCCCCGTCCGGCCCCATGGGGCGGGGCGTGGTTCGACGATGATCCGGCGTGC 660
Q S G P S G P M G R G V V A A M I R R A

661 CCTCGCCGGCGAGCCGCTCACCATGTGGCAGCAGCGGCGGCTGCGCCGCGACCTGTGCA 720
L A G E P L T M W H D G G V R R D L L H

721 CGTCGAGGACGTGGCCACCGGTTCCGCGCGCGCTGGAGCACCACGACGCGCTGGCCGG 780
V E D V A T A F A A A L E H H D A L A G

781 CGGCACGTGGGCGCTGGGCGCCGACCGATCCGAGCCGCTCGGCGACATCTTCCGGGCGGT 840
G T W A L G A D R S E P L G D I F R A V

841 CTCCGGCAGCGTCCGCGGCGAGCCGGCAGCCCCGCGTTCGACGTGGTCACCGTGCCCGC 900
S G S V A R Q T G S P A V D V V T V P A

901 GCCCGAGCACGCCGAGGCCAACGACTTCCGCGAGCAGACATCGACTCCACCGAGTTCCG 960
P E H A E A N D F R S D D I D S T E F R

FIG. 4B-1

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961 CAGCCGGACCGGCTGGCGCCCCGGGTTTCCCTCACCAGCGGCATCGACCGGACGGTGGC 1020
S R T G W R P R V S L T D G I D R T V A

1021 CGCCCTGACCCCCACCGAGGAGCACTAGTCCGGGTACTGCTGACGTCCTTCGCGCACCGC 1080
A L T P T E E H * V R V L L T S F A H R

1081 ACGCACTTCCAGGGACTGGTCCCGCTGGCGTGGGGCGCTGCGCACCGCGGTCACGACGTG 1140
T H F Q G L V P L A W A L R T A G H D V

1141 CGCGTGGCCGCCAGCCCCGCTCACCAGCGCGGTATCGGCGCCGGTCTCACCGCGGTA 1200
R V A A Q P A L T D A V I G A G L T A V

1201 CCCGTCGGCTCCGACCACCGGCTGTTCTGACATCGTCCCGGAAGTCGCCGCTCAGGTGCAC 1260
P V G S D H R L F D I V P E V A A Q V H

1261 CGCTACTCCTTCTACCTGGACTTCTACCAACCGCGAGCAGGAGCTGCACTCGTGGGAGTTC 1320
R Y S F Y L D F Y H R E Q E L H S W E F

1321 CTGCTCGGCATGCAAGAGGCCACCTCGCGGTGGGTATACCCGGTGGTCAACAACGACTCC 1380
L L G M Q E A T S R W V Y P V V N N D S

1381 TTCGTCGCCGAGCTGGTTCGACTTCGCCCGGGACTGGCGTCTGACCTGGTGCTCTGGGAG 1440
F V A E L V D F A R D W R P D L V L W E

1441 CCGTTCACCTTCGCCGGCGCCGTCGCGGCCCGGGCCTGCGGAGCCGCGCACGCCCGGCTG 1500
P F T F A G A V A A R A C G A A H A R L

1501 CTGTGGGGCAGCGACCTCAACGGCTACTTCCGCGGCCGGTTCAGGCGCAACGCCTGCGA 1560
L W G S D L T G Y F R G R F Q A Q R L R

1561 CGGCCGCCGAGGACCGGCCGGACCCGCTGGGCACGTGGTGTGACCGAGGTGCGGGGGCGC 1620
R P P E D R P D P L G T W L T E V A G R

1621 TTCGGCGTCGAATTCGGCGAGGACCTCGCGGTGCGGCAGTGGTTCGGTTCGACCAATTGCCG 1680
F G V E F G E D L A V G Q W S V D Q L P

1681 CCGAGTTTCCGGCTGGACACCGGAATGGAACCGTTGTCCGCGGACCCCTGCCCTACAAC 1740
P S F R L D T G M E T V V A R T L P Y N

1741 GGCGCGTCGGTGGTTCGGACTGGCTCAAGAAGGGCAGTGGGACTCGACGCATCTGCATT 1800
G A S V V P D W L K K G S A T R R I C I

1801 ACCGGAGGGTTCTCCGGACTCGGGCTCGCCGCCGATGCCGATCAGTTCGCGCGGACGCTC 1860
T G G F S G L G L A A D A D Q F A R T L

FIG. 4B-2

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1861 GCGCAGCTCGCGCGATTTCGATGGCGAAATCGTGGTTACGGGTCCGGTCCGGATACCTCC 1920
A Q L A R F D G E I V V T G S G P D T S

1921 GCGGTACCGGACAACATTTCGTTTGGTGGATTTCGTTCCGATGGGCGTTCTGCTCCAGAAC 1980
A V P D N I R L V D F V P M G V L L Q N

1981 TGCGCGGCGATCATCCACCACGGCGGGGCCGGAACCTGGGCCACGGCACTGCACCACGGA 2040
C A A I I H H G G A G T W A T A L H H G

2041 ATTCCGCAAATATCAGTTGCACATGAATGGGATTGCATGCTACGCGGCCAGCAGACCGCG 2100
I P Q I S V A H E W D C M L R G Q Q T A

2101 GAACTGGGCGCGGGAATCTACCTCCGGCCGGACGAGGTCGATGCCGACTCATTGGCGAGC 2160
E L G A G I Y L R P D E V D A D S L A S

2161 GCCCTCACCCAGGTGGTCGAGGACCCACCTACACCGAGAACGCGGTGAAGCTTCGCGAG 2220
A L T Q V V E D P T Y T E N A V K L R E

2221 GAGGCGCTGTCCGACCCGACGCCGCGAGGATCGTCCCGCGACTGGAGGAACTCACGCGC 2280
E A L S D P T P Q E I V P R L E E L T R

2281 CGCCACGCGCGCTAGCGGTTTCCGACCGACAAGTCCGTCCGACAGCACACCTCCGGAGGG 2340
R H A G *

2341 AGCAGGGATGTACGAGGGCGGGTTCGCCGAGCTTTACGACCGGTTCTACCGCGCCGGGG 2400
M Y E G G F A E L Y D R F Y R G R G

2401 CAAGGACTACGCGGCCGAGGCCGCGCAGGTTCGCGCGGCTGGTCAGAGACCGCTGCCCTC 2460
K D Y A A E A A Q V A R L V R D R L P S

2461 GGCTTCCTCGCTGCTCGACGTGGCTGCGGGACCGGCACCCACCTGCGCCGGTTCGCCGA 2520
A S S L L D V A C G T G T H L R R F A D

2521 CCTCTTCGACGACGTGACCGGGCTGGAGCTGTGCGCGCGATGATCGAGGTTCGCCGGCC 2580
L F D D V T G L E L S A A M I E V A R P

2581 GCAGCTCGGCGGCATCCCGGTGCTGCAGGGCGACATGCGCGACTTCGCGCTGGATCGCGA 2640
Q L G G I P V L Q G D M R D F A L D R E

2641 GTTCGACGCCGTACCTGCATGTTTCAGCTCCATCGGGCACATGCGCGACGGCGCCGAGCT 2700
F D A V T C M F S S I G H M R D G A E L

2701 GGACCAGGCGCTGGCGTCCCTTCGCCCCCACCCTCGCCCCGGCGGCGTCTGGTGGTTCGA 2760
D Q A L A S F A R H L A P G G V V V V E

2761 ACCGTGGTGGTTCCCGGAGGACTTCCTCGACGGCTACGTGGCCGGTGACGTGGTGC CGCA 2820
P W W F P E D F L D G Y V A G D V V R D

FIG. 4B-3

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2821 CGGCGACCTGACGATCTCGCGCTCTCGCACTCCGTGCGCGCCGGCGGCGCGACCCGGAT 2880
G D L T I S R V S H S V R A G G A T R M

2881 GGAGATCCACTGGGTGCTGGCCGACGCGGTGAACGGTCCGCGGCACCACGTGGAGCACTA 2940
E I H W V V A D A V N G P R H H V E H Y

2941 CGAGATCACGCTCTTCGAGCGGCAGCAGTACGAGAAGGCCCTTCACCGCGGCCGGTTGCGC 3000
E I T L F E R Q Q Y E K A F T A A G C A

3001 TGTGCAGTACCTGGAGGGCGGACCCTCCGGACGCGGGTTGTTGTCGGTGTGCGCGGATG 3060
V Q Y L E G G P S G R G L F V G V R G *

3061 ACCCGTGCCTCGCGTTTTCCGTTCTGGCAGAGGTGATCCGCTCCACGGGCCCTTTCCCC 3120

3121 GCCGTGACCGGACCCTTACAGTGAGTGCGGGTCTTGATCGACAACGCCCGGCGGCAGCAA 3180

3181 GCGGAGCCGTCGACGACACCGCAGGGAGAGTCGATGGGTGATCGGACCGGCGACCGGACG 3240
M G D R T G D R T

3241 ATTCCGGAATCCTCGCAGACCGCAACGCGTTTCTGCTCGGCGACGGCGGAATCCCCACC 3300
I P E S S Q T A T R F L L G D G G I P T

3301 GCCACGGCGGAAACCCACGACTGGCTGACCCGCAACGGCGCCGAGCAGCGGCTCGAGGTG 3360
A T A E T H D W L T R N G A E Q R L E V

3361 GCGCGCGTGCCGTTTCAGCGCCATGGACCGCTGGTTCGTTCCAGCCCGAGGACGGCAGGCTC 3420
A R V P F S A M D R W S F Q P E D G R L

3421 GCCACGAGTCCGGGCGCTTCTTCTCCATCGAGGGCCTGCACGTGCGGACGAACCTTCGGC 3480
A H E S G R F F S I E G L H V R T N F G

3481 TGGCGGCGGGAAGTGGATCCAGCCCATCATCGTGCAGCCCGAGATCGGCTTCTCGGCCTC 3540
W R R D W I Q P I I V Q P E I G F L G L

3541 ATCGTCAAGGAGTTTCAGCGGTGTGCTGCACGTGCTGGCGCAGGCCAAGGCCGAGCCGGG 3600
I V K E F D G V L H V L A Q A K A E P G

3601 AACATCAACGCCGTCCAGCTCTCCCCGACCCTGCAGGCGACCCGCGAGCAACTACACCGGC 3660
N I N A V Q L S P T L Q A T R S N Y T G

3661 GTCCACCGCGGCTCGAAGGTCCGGTTCATCGAGTACTTCAACGGCACGCGCCCGAGCCGG 3720
V H R G S K V R F I E Y F N G T R P S R

3721 ATCCTCGTCGACGTGCTCCAGTCCGAGCAGGGCGCGTGGTTCTGCGCAAGCGCAACCGG 3780
I L V D V L Q S E Q G A W F L R K R N R

FIG. 4B-4

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3781 AACATGGTCGTCGAGGTGTTTCGACGACCTGCCCGAGCACCCGAACCTCCGGTGGCTGACC 3840
N M V V E V F D D L P E H P N F R W L T

3841 GTCGCGCAGCTGCGGGCGATGCTGCACCACGACAACGTGGTGAACATGGACCTGCGCACC 3900
V A Q L R A M L H H D N V V N M D L R T

3901 GTGCTGGCCTGCGTCCCGACCCGCTGGAGCGGGACCGGGCCGACGCTGCTGCGCGCG 3960
V L A C V P T A V E R D R A D D V L A R

3961 CTGCCCCGAGGGCTCGTTCAGGCCCGGCTGCTGCACTCGTTCATCGGCGCGGGCACCCCG 4020
L P E G S F Q A R L L H S F I G A G T P

4021 GCCAACAACTGAACAGCCTGCTGAGCTGGATCTCCGACGTGCGCGCCAGGCGCGAGTTC 4080
A N N M N S L L S W I S D V R A R R E F

4081 GTGCAGCGCGGCCCGCCGCTGCCCGACATCGAGCGCAGCGGGTGGATCCGCCGCGACGAC 4140
V Q R G R P L P D I E R S G W I R R D D

4141 GGCATCGAGCACGAGGAGAAGAAGTACTTCGACGTCTTCGGCGTCACGGTGGCGACCAGC 4200
G I E H E E K K Y F D V F G V T V A T S

4201 GACCGCGAGGTCAACTCGTGGATGCAGCCGCTGCTCTCGCCCGCCAACAACGGCCTGCTC 4260
D R E V N S W M Q P L L S P A N N G L L

4261 GCCCTGCTGGTCAAGGACATCGGCGGCACGTTGCACGCGCTCGTGCAGCTGCGCACCGAG 4320
A L L V K D I G G T L H A L V Q L R T E

4321 GCGGGCGGGATGGACGTGCGCCGAGCTGGCGCCTACGGTGCCTGCGCAGCCGACAACTAC 4380
A G G M D V A E L A P T V H C Q P D N Y

4381 GCCGACGCGCCCGAGGAGTTCGACCGGCCCTATGTGGACTACGTGTTGAACGTGCCGCGC 4440
A D A P E E F R P A Y V D Y V L N V P R

4441 TCGCAGGTCCGCTACGACGCATGGCACTCCGAGGAGGGCGCCGGTTCTACCGCAACGAG 4500
S Q V R Y D A W H S E E G G R F Y R N E

4501 AACCGGTACATGCTGATCGAGGTGCCCGCCGACTTCGACGCCAGTGCCGCTCCCGACCAC 4560
N R Y M L I E V P A D F D A S A A P D H

4561 CGGTGGATGACCTTCGACCAGATCACCTACCTGCTCGGGCACAGCCACTACGTCAACATC 4620
R W M T F D Q I T Y L L G H S H Y V N I

4621 CACGTGCGCAGCATCATCGCGTGGCGCTCGGCCGTCTACACCAGGACCGCCGGATGAAAC 4680
H V R S I I A C A S A V Y T R T A G *
M K R

4681 GCGCGCTGACCGACCTGGCGATCTTCGGCGGCCCGAGGCATTCTGCAACCCCTCTACG 4740
A L T D L A I F G G P E A F L H T L Y V

FIG. 4B-5

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4741 TGGGCAGGCCGACCGTCGGGGACCGGGAGCGGTTCTTCGCCCCCCTGGAGTGGGCGCTGA 4800
G R P T V G D R E R F F A R L E W A L N

4801 ACAACAACCTGGCTGACCAACGGCGGACCACTGGTGCGCGAGTTCGAGGGCCGGGTCGCCG 4860
N N W L T N G G P L V R E F E G R V A D

4861 ACCTGGCGGGTGTCCGCCACTGCGTGCCACCTGCAACGCGACGGTCGCGCTGCAACTGG 4920
L A G V R H C V A T C N A T V A L Q L V

4921 TGCTGCGCGCGAGCGACGTGTCCGGCGAGGTCGTCATGCCCTTCGATGACGTTCCGGGCCA 4980
L R A S D V S G E V V M P S M T F A A T

4981 CCGCGCACGCGGCGAGCTGGCTGGGGCTGGAACCGGTGTTCTGCGACGTGGACCCCGAGA 5040
A H A A S W L G L E P V F C D V D P E T

5041 CCGGCTGCTCGACCCGAGCACGTGCGCTGCTGGTCAACCGCGACGGGCGCGATCA 5100
G L L D P E H V A S L V T P R T G A I I

5101 TCGGCGTGCACTCTGCGGCAGGCCCCGCTCCGGTCGAGGCGCTGGAGAAGATCGCCGCCG 5160
G V H L W G R P A P V E A L E K I A A E

5161 AGCACCAGGTCAAACCTCTTCTTCGACGCCGCGCACGCGCTGGGCTGCACCGCCGGCGGGC 5220
H Q V K L F F D A A H A L G C T A G G R

5221 GGCCGGTCGGCGCCTTCGGCAACGCCGAGGTGTTGAGCTTCCACGCCACGAAGGCGGTCA 5280
P V G A F G N A E V F S F H A T K A V T

5281 CCTCGTTCGAGGGCGGCGCCATCGTCACCGACGACGGGCTGCTGGCCGACCGCATCCGCG 5340
S F E G G A I V T D D G L L A D R I R A

5341 CCATGCACAACCTTCGGGATCGCACCGGACAAGCTGGTGACCGATGTCGGCACCAACGGCA 5400
M H N F G I A P D K L V T D V G T N G K

5401 AGATGAGCGAGTGCGCCGCGGCGATGGGCCTCACCTCGCTCGACGCCTTCGCCGAGACCA 5460
M S E C A A A M G L T S L D A F A E T R

5461 GGGTGCACAACCGCCTCAACACGCGCTCTACTCCGACGAGCTCCGCGACGTGCGCGGCA 5520
V H N R L N H A L Y S D E L R D V R G I

5521 TATCCGTGCACGCGTTCGATCCTGGCGAGCAGAACAACCTACAGTACGTGATCATCTCGG 5580
S V H A F D P G E Q N N Y Q Y V I I S V

5581 TGGACTCCGCGGCCACCGGCATCGACCGCGACCACTTGCAAGGCGATCCTGCGAGCGGAGA 5640
D S A A T G I D R D Q L Q A I L R A E K

5641 AGGTTGTGGCACAACCTACTTCTCCCCGGGTGCCACCAAGATGCAGCCGTACCGGACCG 5700
V V A Q P Y F S P G C H Q M Q P Y R T E

FIG. 4B-6

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5701 AGCCGCCGCTGCGGCTGGAGAACACCGAACAGCTCTCCGACCGGGTGCTCGCGCTGCCCA 5760
P P L R L E N T E Q L S D R V L A L P T

5761 CCGGCCCCGCGGTGTCCAGCGAGGACATCCGGCGGGTGTCGACATCATCCGGCTCGCCG 5820
G P A V S S E D I R R V C D I I R L A A

5821 CCACCAGCGGCGAGCTGATCAACGCGCAATGGGACCAGAGGACGCGCAACGGTTCGTGAC 5880
T S G E L I N A Q W D Q R T R N G S *

5881 GACCTGCGCCACAAGTGCCAGGAGGTTTCGCTCCCCGATGAACACAACCTCGTACGGCAACC 5940
M N T T R T A T

5941 GCCCAGGAAGCGGGGGTCGCGGACGCGGCGCGCCCGGACGTCGACCGGCGGGCGGTCTGTG 6000
A Q E A G V A D A A R P D V D R R A V V

6001 CGGGCGCTGAGCTCGGAGGTCTCCCGCGTCACCGGCGCGGTGACGGTGACGCCCACGTG 6060
R A L S S E V S R V T G A G D G D A H V

6061 CAGGCGCCCGGCTCGCCGACCTCGCCGCGCACTACGGGGCGCACCCGTTACGCGCGCTG 6120
Q A A R L A D L A A H Y G A H P F T P L

6121 GAGCAGACGCGTGCGCGGCTCGGCCTGGACCGCGCGGAGTTCGCCCACCTGCTCGACCTG 6180
E Q T R A R L G L D R A E F A H L L D L

6181 TTCGGCCGCTCCCGGACCTGGGCACCGCGGTGGAGCACGGTCCGGCGGGCAAGTACTGG 6240
F G R I P D L G T A V E H G P A G K Y W

6241 TCCAACACGATCAAGCCGCTGGACGCCGCGAGGCGCACTGGACGCGGCGGTCTACCGCAAG 6300
S N T I K P L D A A G A L D A A V Y R K

6301 CCTGCCTTCCCCTACAGCGTCGGCCTGTACCCCGGGCCGACGTGCATGTTCCGCTGCCAC 6360
P A F P Y S V G L Y P G P T C M F R C H

6361 TTCTGCGTGCGGGTGACCGGTGCCCGCTACGAGGCCGATCGGTCCCGGCGGGCAACGAG 6420
F C V R V T G A R Y E A A S V P A G N E

6421 ACGCTGGCCGCGATCATCGACGAGGTGCCACGGACAACCGAAGGCGATGTACATGTG 6480
T L A A I I D E V P T D N P K A M Y M S

6481 GGCGGGCTCGAGCCGCTGACCAACCCCGGTCTCGGCGAGCTGGTGTGCGACGCGCGCGGG 6540
G G L E P L T N P G L G E L V S H A A G

6541 CGCGGTTTCGACCTCACCCTCTACACCAACGCCTTCGCCCTCACCAGCAGACGCTGAAC 6600
R G F D L T V Y T N A F A L T E Q T L N

6601 CGCCAGCCCGCCTGTGGGAGCTGGGCGCGATCCGCACGTCCCTCTACGGGCTGAACAAC 6660
R Q P G L W E L G A I R T S L Y G L N N

FIG. 4B-7

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6661 GACGAGTACGAGACGAACACCGGCAAGCGGGCGCTTTTGAACGCGTCAAGAAGAACCTG 6720
D E Y E T T T G K R G A F E R V K K N L

6721 CAGGGCTTCCTGCGGATGCGCGCCGAGCGGGACGCGCCGATCCGGCTCGGCTTCAACCAC 6780
Q G F L R M R A E R D A P I R L G F N H

6781 ATCATCTGCGGGACGGGCGGACCGGCTCACCGACCTCGTTCGACTTCATCGCCGAGCTC 6840
I I L P G R A D R L T D L V D F I A E L

6841 AACGAGTCCAGCCCGCAACGGCCGCTGGACTTCGTGACGGTGCGCGAGGACTACAGCGGC 6900
N E S S P Q R P L D F V T V R E D Y S G

6901 CGCGACGACGGCCGGCTGTCTGGACTCCGAGCGCAACGAGCTGCGCGAGGGCCTGGTGCGG 6960
R D D G R L S D S E R N E L R E G L V R

6961 TTCGTCGACTACGCCGCCGAGCGGACCCCGGGCATGCACATCGACCTGGGCTACGCCCTG 7020
F V D Y A A E R T P G M H I D L G Y A L

7021 GAGAGCCTGCGGGCGGGTGTGGACGCCGAGCTGCTGCGCATCCGGCCGGAGACGATGCGT 7080
E S L R R G V D A E L L R I R P E T M R

7081 CCCACCGCGCACCCCCAGGTCCGGGTGCAGATCGACCTGCTCGGCGACGTCTACCTCTAC 7140
P T A H P Q V A V Q I D L L G D V Y L Y

7141 CGCGAGGCGGGCTTCCCGGAGCTGGAGGGCGCCACCCGCTACATCGCGGGCCGGGTCAAC 7200
R E A G F P E L E G A T R Y I A G R V T

7201 CCGTCGACCAGCCTGCGCGAGGTGGTGGAGAACTTCGTGCTGGAGAACGAGGGCGTGCA 7260
P S T S L R E V V E N F V L E N E G V Q

7261 CCCCCCCCCGCGACGAGTACTTCTCGACGGCTTCGACCAGTCGGTGACCGCACGGCTC 7320
P R P G D E Y F L D G F D Q S V T A R L

7321 AACCAGCTCGAACGAGACATCGCCGACGGGTGGGAGGACCACCGGGCTTCTGCGCGGA 7380
N Q L E R D I A D G W E D H R G F L R G

7381 AGGTGAACCGGAGTTGCGAGTACGTGAGCTGGCGGTGGCGGGCGGTTTCGAGTTCACCC 7440
R * V A G G F E F T P

7441 CGACCCGAAGCAGGACCGGCGGGGCTGTTCGTGTCTCCGCTGCAGGACGAGGCGTTCGT 7500
D P K Q D R R G L F V S P L Q D E A F V

7501 GGGCGCGGTGGGCCATCGGTTCCCCGTCGCCCAGATGAACCACATCGTCTCCGCCCGGG 7560
G A V G H R F P V A Q M N H I V S A R G

7561 CGTGCTGCGCGGGCTGCACTTCACCACCACCCCGCGGGGCGAGTGCAAGTACGTCTACTG 7620
V L R G L H F T T T P P G Q C K Y V Y C

FIG. 4B-8

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7621 CGCGCGCGGCGGGCGCTCGACGTCATCGTCGACATCCGGGTCGGCTCGCCGACGTTCCG 7680
A R G R A L D V I V D I R V G S P T F G

7681 GAAGTGGGACGCGGTGGAGATGGACACCGAGCACTCCGGGCGGTCTACTTCCCCAGGGG 7740
K W D A V E M D T E H F R A V Y F P R G

7741 CACCGCGCACGCCTTCCTCGCGCTTGAGGACGACACCCTGATGTCGTACCTGGTCAGCAC 7800
T A H A F L A L E D D T L M S Y L V S T

7801 GCCGTACGTGGCCGAGTACGAGCAGGCGATCGACCCGTTGACCCCGCGCTGGGTCTGCC 7860
P Y V A E Y E Q A I D P F D P A L G L P

7861 GTGGCCCGCGGACCTGGAGGTCGTGCTCTCCGACCGGACACGGTGGCCGTGGACCTGGA 7920
W P A D L E V V L S D R D T V A V D L E

7921 GACCGCCAGGCGGCGAGGGATGCTGCCCCACTACGCCGACTGCCTCGGCGAGGAGCCCGC 7980
T A R R R G M L P D Y A D C L G E E P A

7981 CAGCACCGGCAGGTGACGGGTCCCGAGCACGATCTGTTCGAAGTGGCGCAGGCGCTCGTC 8040
S T G R *

8041 GTCGCGGTCGA 8051

FIG. 4B-9

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```

1          vngidsprq  litllGaSGf  vGsavlreLr  dhpv.rlrav  sRggapavpp  50
eryBIV      ascF      .llitGvSGy  iGshlvnyLa  nlgyeyiygi  sRneildqdi
          rfbJ      ...mtflke  yvivsGaSGf  iGkhllleaLk  k.sgisvvai  tRdvi...knn
          strL      ...mspyprp  rwlvtGaSGm  lGreltpLLd  rrga.avtal  gRghl.ditd
Consensus  -----G-SG- -G-----L- -R-----

```

```

51          gaaevedlra  dllepgraaa  aiedadvivh  lvahaAggst  wrsatsdpe.  100
eryBIV      ascF      nql.llniki  fqldrdslpd  ilkrvrpdv.  .vihlAscfl  sqhsyknike
          rfbJ      sna.lanvrw  cswdniellv  eelsidsali  giihlAtey.  .ghktsslin
          strL      gaa.....  ....vrsav  aehrpaavvn  caawtAvd..  ..eaesepal
Consensus  -----A-----

```

```

101         aervNvglmh  dLvgalhrr  rstppvllly  staqaanpsa  asryaqqkte  150
eryBIV      ascF      iiksNvefpt  eLlea.....  ....mndv  gvkkiintgt  swqcfnsdty
          rfbJ      iedaNvikpl  kLldl.....  ....aiky  radiflntd.  sffakkdfnv
          strL      amavNgegpr  hLaqa....c  ravgavllql  stdyvfpgsg  grpyredhpt
Consensus  ----N-----L-----

```

FIG. 5A-1

	151		200
eryBIV	aerilrkatd	egrvrgvilr lpa.....	..vyqsgps gpmgrgvvaa
ascF	npvnlyaask	qafedilkfy inaeafsain	lklfdtyGgv dkrrklis.l
rfbJ	qhmrpyiitk	rhfdeighyy anmhdisfvn	mrlehvyGpg dgenkfipyi
strL	gprrtvygctk	rageravlev lpdtgyivrt	awlygagGp.nfvak
Consensus	-----	-----	-----G-----
	201		250
eryBIV	mirralagep	ltmwhdggvr rd1.lhveDv	atAFaaaleh hdalag...g
ascF	lddiaknnkq	ldmspgeql1 d..lvhinDv	crAFkiaidk lcelpseyvv
rfbJ	idclnkkqsc	vkcttgeqir d..fifvdDv	vnAYltilen rkevps...yt
strL	mirleadedt	vlvddqhgq ptwtadlaDr	laALgaaala gtapagiyha
Consensus	-----	-----D-----	-----AF-----

FIG. 5A-2

	251		300
eryBIV	twalgadrse	plgDifravs	gsva..rqtg
ascF	sygvsnkyrv	tlkElvsiye	..rv...nnv
rfbJ	eyqvgtgagv	slkDflvylq	ntmm...pgs
strL	tntggttwna	lapEtfrllg	adparvrptt
Consensus	-----	-----	-----
			spavdvvtvp
			apehaeandf
			yrnrevmvp
			qrdneimfsv
			.prysvldqs

	301		345
eryBIV	rsddidstef	rsrtgwrprv	sltdgidrtv
ascF	.tniqnl...	...pdwevvi	plsqglky..
rfbJ	.annknlkam	gwkpndydkk	gieellkrl.
strL	rwkaaglepl	rhw.....ra	altesf....
Consensus	-----	-----	-----
			palcgragrp
			vpgpr

FIG. 5A-3

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```

1      eryBVII  .... 50
      strM.    ..   .vrplsVqga wlsetrafad dRGefqelys arslrgalGy
      rfbC     .....vm iviktaIpdv lilepkvfgD eRGfFfesYn qqtFeeliGr
      rfbF     .....m  kctklsIpev ilfeprifed dRGhFfesFn lakFqesiGr
      ascE     lgvivphyIm ifkkldIegc yIiefnkfid sRGtFvktFh sdfFsen.Gi
      Consensus -----I---D-----RG-F-----F-----G-

```

```

51      eryBVII 100
      strM     rfpvaqmnhI vSargVlRGl HFtttppgqc KyVycarGra LDVivDiRvg
      rfbC     dpgvaqvnrs vSrrgVlRGv HFaqIppsqA KyVtclsGav LDVvvDiRtg
      rfbF     kvtfvqdnhs kSkknVlRGl HFqrgenagg KlVrcavGev FDVavDiRke
      ascE     qvtfvqsnes ySkqnViRGl HYq.virpqq KlVrvveGev FDIavDlRks
      Consensus vldmreefys iSaknViRGm HFqmpaehd KlVycvnGav LDViIdiRkd
      -----S---V-RG--HF-----K-V-----G--LDV--D-R--

```

```

101      eryBVII 150
      strM     SpTFgkwdaV eMdtehfrav YfprGtaHaF laLeddltlms Ylvstpyvae
      rfbC     SpTYraWeaV rLddph.rsl YveaGlghSF maLddavvv Yltsqgyaag
      rfbF     SpTFgqWvgV nLsaenkrql WipeGfaHgF vtLseyaefl Ykatnyysps
      ascE     SpTFgqWvgV lLsdknqhql WipeGfgHgF qvLspsakfq YmvtDywyype
      Consensus SkTYgeYfsI eLsyenslal WvpkGlaHgF lsLadnsimf Yktssvhnve
      S-TF--W--V -L-----W---G--H-F--L-----Y-----

```

FIG. 5B-1

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eryBVII	151	yEqalDpfdp	alglpwpadl	evvlsDrdtv	avdletarr	gmldpyadcl	200
strM		rEhgVhpldp	dlgiawpdgi	epvlsEkdrq	apgiaemerr	gllpdyeecl	
rfbC		sEgsIlwnde	aigiewp...	fsqlpE...	...lsakdaa	aplldqallt	
rfbF		hDrcIrfnds	dinikwk...	egiisEqqvi	eyklsskdis	gnsladaevf	
asCE		cDsgIkwnsf	gfkwpid...	npiisEkdns	lcyfdefdss	f.....	
Consensus		-E--I-----	-----E-----	-----E-----	-----	-----	
eryBVII	201	geepastgr	..				
strM		afrrslcerg	tg				
rfbC		e.....	..				
rfbF					
asCE					
Consensus		-----	---				

FIG. 5B-2

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FIG. 5C-1

1	50
eryCIVmk raltDLaifg gpeaFlhtly vgrptVgd..rerFfa
eryCImd vpfldLqa.. ayleLrsdid qACrrVlg..sgwY..
ascC	...msqeelr qqiaelVaq. aetaMapkpf eAgksVvpps gkvigtkelq
dnrJ vstyvWqyln eyreeradiI dAVetVfe..sgqL..
prgI
strSmssFqelp rwpqLtdddi eAavaalr..snrL..
Consensus	-----L-----L----- -A---V-----
51	100
eryCIV	rlewalnnnw Ltnggplvre FEgrvAdL.. aGvrHcVatc natvAlqLvL
eryCI Lhgpe..nea FEaeFAaY.. cenaHcVtvg SGcdAleLsL
ascC	lmveasldgw L.ttggrfnda FEklgeYl. .GVpyvlttt SGssAnlLAL
dnrJ ilgts..vrs FEeeFAaY.. hGlpyctgvd nGtnAlvLgL
prgI,sgp..igq LEaeFlaFlD hGvryaVtfn SGtsAlIaAY
strS vgggnstvee FEaaLaa..g qGveHaVavs tGtaaVhLAL
Consensus	-----L----- FE---A-Y-- -G--H-V--- SG--A--LAL
101	150
eryCIV	rAs..... .dv..sgEVV mPsmTFaata haaswlGleP VFcDVDpeTg
eryCI	vAL..... .gVgqGDEVI vPshTFiaTw lgV.pvGAVP VpVEpEgvsh
ascC	tALtspklgv ralkPGDEVI tvaagFptTv nptiqnGlip VFVDVDipT.
dnrJ	rAL..... .gIgPGDEVI tvsnTaaPTv vaIdavGATP VFVDVhe...
prgI	fAL..... .gVreGvEaa gPaLYhaal spVfalrgdv VLVdIDpvsr
strS	hAL..... .dVgPGDEVI vPthTFigsa spVtylGARp VFADvtpdTh
Consensus	-AL----- --V-PGDEVI -P--TF--T- --V---GA-P VFVDVD--T-

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FIG. 5C-2

	151		200
eryCIVLDP	ehVaslVTpr	TgAligVHlW GrpapvEale kIAaeHqvkl
eryCI	t.....LDP	alVEqAITpr	TaAilpVHlY GhpaDLdalr aIAdrHgLaI
ascCynvna	sliEAAVsDk	TkAlmiaHtI GnlfDLaeVr rVAdkynLwI
dnrJ	...enylMDt	grlrsVlgpr	TrcllpVHlY GqsvDMtpVl elAaeHdLkV
prgI	g.....LDP	kaIEAAITEn	TrvVtvVHqW GhpcDMDaIl gVAerygLrV
strS	c.....LDP	dsvksllIGer	TkAIvvVHIn GiaadMaaIt aVAaeagvpV
Consensus	-----LDP	--VEAAITE-	T-AI--VH-Y G---DMD-V- -IA--H-L-V
	250		
eryCIV	201	ffDaAhAlGc	tagGrpVGaF GnaevFS.Fh atKavtsf.E GGAIVTdDgll
eryCI		VEDvAQAVGa	rhrGhrVGag snaaafs.FY PgKnlgAlGD GGAVVtDpal
ascC		IEDccdaLGS	tydGkmaGtF GdigitVS.FY Pahhitm.GE GGAVfTqsaEL
dnrJ		IEDcAQAhGa	rrhGrLVGtq Ghaaafs.FY PtKvlgAyGD GGAVVtpDaev
prgI		IEDcshAhGS	rYkGkvpGtF GdaavFS.Lq anKavyA.GE GGilVTdDalv
strS		IEDaAQAlgt	eigGrpIGgF GdlacvSlFF eqKvitsgGE GGAVITdnpvY
Consensus		IED-AQA-G-	-Y-G--VG-F G-----FS-FY P-K---A-GE GGAVVT-D--L
	300		
eryCIV	251	aDRiRaMhn	FGiapdk... ..lvtDvG
eryCI		aERiRlLRn	YG.....sk.qk.YvhvvrG
ascC		ksiiiesFRd	WGrdcycapG cdntckkrfg qqlgslpfgY dhkYtyshlG
dnrJ		drRlRrLRy	YG.....mg.eryYvvdtpG
prgI		qDRatlLat	tG.....t..vpg
strS		aERvRrLRs	hGegpvsgs.pgmiwahevG
Consensus		-ER-R-LR-	YG-----Y-----G

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```

301          tNgkMsEcaA AmgltsLdaF aetrvhnrln halysdeLrd vrGisvhafd
eryCIV      tNaRLdELQA AvlrvkLrhL DdWnarRttl aghyqteLkd vpGitlpeth
eryCI       yNikitDMQA AcglaqLepi EeFvekRkan fkyLkdalqs cadf.ielpe
ascC        hNsRLdEvQA eilrrkLrrL DaYvegRrav arryeegLgd ldGlvlppt..
dnrJ        lghRr.....
prg1        yNvRLtsvQA psaspsnkrl gdlveaRrrn aaylserLag veGlelpvpep
strS        -N-RL-E-QA A-----L--L D-W---R--- -L- -G-----
Consensus

```

```

351          pgeqnnyqYv iisvdsaatg idr..dqlqa iLraekVvaq pyFspgCHqm
eryCIV      pwads..aWh lFvlrcenrD .....hLqr hLtdagVqtl ihYptpvHls
eryCI       atensdpsWf gFpI.tlkeD sgvsridLvK fLdeakVgtr llFagnltrq
ascC        iaegn dhvYy vYvVrhperD .....rile aLtaydiHln isYpwpvHtm
dnrJ        .....
prg1        p...gtthaYw kYaVrvvpGD grrsaadiaa hLrsrgVpvl lrYpyplHkq
strS        -----W- -Y-----D -----L-- -L-----V--- --Y-----H--
Consensus

```

FIG. 5C-3

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```

401          qpYrte...p plrLentEqL sdrvLaLPtg PavssEdirr Vcdiirlaat
eryCIV          paYAdlgl.p pGsfPvaESL agevLsLPig PhLsrEaaDh VIatlkaga
eryCI          pyFhdvkyRv vGeLtnDRI mnqtFwigiy PgLthDhLDY Wskfeefg
ascC          sgFAhlg.Yg pGdLpvtERL ageiFsLPmy PsLrpDaqEk Vidavrevvg
dnrJ          .....
prg1          paFAe...Yh gvslpvaERL sqellaLPsh PgLveghlDh aVeevrkava
strS          --FA----Y- -G-L---ERL ----L-LP-- P-L--E--D- VW-----
Consensus

```

```

451          sgelinaqwd qtrngs
eryCIV          .....
eryCI          lnf.....
ascC          sl.....
dnrJ          .....
prg1          .....
strS          s.....
Consensus          -----

```

FIG. 5C-4

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```

1      50
    eryBV      vrVlltsfah rthFqGlvPL AWALRtAGHD VRVaaqPALT DaVigAGLTA
    eryCIII     mrVvfssmAs kshLfGlvPL AWAFraAGHE VRVvasPALT EditaaGLTA
    dnrS      mkVlvtafam daHFngvVPL AWALRaAGHD VRVasqPALT DsItrAGLTA
Consensus  --V-----A- --HF-G-VPL AWALR-AGHD VRV---PALT D-I--AGLTA

51      100
    eryBV      VPVGsDhrIf divpeaaqV hrysfyldFy hregelhsWE fLlgmqeats
    eryCIII     VPVGtDvdLv dfmthaghdI idyvrsldFs erdpatltwE hLrgmqtvlt
    dnrS      VPVGtDhqVq aamgamapGV falhlnpDYl enrpellldE fLeastsmLt
Consensus  VPVG-D-----V -----DF- -----E -L-----

101     150
    eryBV      rwvYpvnnd sFvaeLvDfa rdWRPDLVlW EPFTFagaVA aracGaahAR
    eryCIII     ptfYalmspd tLlegMVsfC rkWRPDLViW EPLTFaapIA gavtGtphAR
    dnrS      aafYaqinnD sMideMVdFa awWRPDLVvW EPFTFggava aqvTGaaqAR
Consensus  ---Y-----D --I--MV-F- --WRPDLV-W EPFTF---VA ---G---AR

```

FIG. 5D-1

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151				200
eryBV	LLWGsDltgy	frgrFqaqrl	rrPpEdRpdP	LgtWLTevag rfGv.eFgeD
eryCIII	LLWGpDittR	arqnFlgllp	dqPeEhRegp	LaeWLTwtle kyGgpaFdeE
dnrS	LLWGpDlflr	vhdrFqqvlh	evPaErRdda	LeeWLTwtle rhGa.aFgpe
Consensus	LLWG-D----	----F-----	--P-E-R---	L--WLT----- --G---F--E
201				250
eryBV	lavGqWsvDq	lPpsfRLdtg	meTVvaRtlp	YNG...asVVP dWLkksatr
eryCIII	vVvGqWtIDp	aPaaIRLdtg	lKTVgmRyvD	YNG..psVVP eWLhdeper
dnrS	visGhWtIDq	mPpsvRFaTa	rpTVpmRfvp	YNGpvpaVVP pWLradpgrp
Consensus	---G-W-ID-	-P----RL-T-	--TV--R---	YNG-----VVP -WL-----
251				300
eryBV	RIciTgGfsg	lgla.adadq	fartLaqlar	fDgEIVvTgs gpdtSav...
eryCIII	RVclTlGiss	rensiggvs.	ieellgavgd	vDaEIIaTfd aqlegvani
dnrS	RVllTqGite	rstgftglpr	agellasiae	lDaEVVaTvk aeereg1pp1
Consensus	RV--T-G---	-----	-----L-----	-D-EIV-T-- -----

FIG. 5D-2

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301
 eryBV PdNIRlVdfv pMgvllqnCA AiIHhGGaGt WaTAhHGIP QisvahewDc 350
 eryCIII PhNVRtVgfv pMhallptCA AtVHHGgpGs WhTAaiHGVP QvilpdgWdt
 dnrS PGNVRvVds1 sLhvvLpsCA AvVHHGgGt WaTAalHGVP QlalaWqWdd
 Consensus P-NVR-V--- -M---L---CA A-VHHGG-G- W-TA--HGVP Q-----WD-

351
 eryBV mlRgqqtaeL GAGiYlrp.. devdadslas altqVvedPt ytenAvklRe 400
 eryCIII gvRaqrqtqef GAGIaLp..v peltpdqlre svkrVldDpa hragAarmRd
 dnrS vFRagglekL GAGIfLpphg egasagrVrd rlaqVlaEPs frqgAariRa
 Consensus --R-----L GAGI-L-----V---DP- ----A---R-

401
 eryBV EaLsdptPqe IVprlEeLtr rhag..... 433
 eryCIII DmLaePsPae VVgicEeLaa grrepr....
 dnrS EmLrtPaPga VVptlEqLta rhrapagggv rh
 Consensus E-L--P-P-- VV---E-L-- ----

FIG. 5D-3

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1
 50
 eryCVI MYegg.fAel YDrfyrgRgK DYaaeaqva rlvrdrlpsA ssLLDVACGT
 srmX MYendsaAev YDllyqdr.K DYageaarvt dliirertpdA asLLDIACGT
 rdmD MYgad.lArv YDlvhreRgK DYrardrggr rrgpaeqagA grLLDVACGT
 Consensus MY-----A-- YD-----R-K DY-----A --LLDVACGT

51
 100
 eryCVI GtHLrrFAdL FddVtGLELS aaMievArpQ LgGlpvlqGD MRdFaIdref
 srmX GtHLeaFAkL YdrVsGLELS ewMaarAeer LpGVtlhrGD MRaFdLgetF
 rdmD GgHLrhFAdL FahVeGvELS epMaeeAraa LpGVtvhaGD MRdFrLgttF
 Consensus G-HL--FA-L F--V-G-ELS --M---A--- L-GV-----GD MR-F-L---F

101
 150
 eryCVI DaVtCMFSSI GhMrdgAELd qAlasfARHL apgGVvvVEP WWFpEdFIDG
 srmX DaVvCMFSSI GyLettADLe dAvaamARHL tadGVlaVEP WYFpDtFIDG
 rdmD DvVtCMFgSV GyMtsvAELg rAlrmfARHL epgGVavVDP WWFyEtFaDG
 Consensus D-V-CMF-SI G-M---AEL- -A-----ARHL ---GV---VEP WWF-E-F-DG

FIG. 5E-1

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eryCVI	151	yVagdvvrD.	.gdltIsRVS	HsvRaGgatr	MEIHWvVAda	vnGprHhvEh	200
srnX		hVsthalrta	pgdqgVaRVS	HStReGgrtr	MEIHylIAht	aeGirHrsEv	
rdmD		hVsadivtv.	.dgvTVsRVS	HSaRrGrtsh	MDVHFvVAep	gaGaQHfvDt	
Consensus		-V-----	-----V-RVS	HS-R-G----	MEIH--VA--	--G--H--E--	
eryCVI	201	yeitLFeRqq	YEKAftaAGc	avqYleggps	grGLFvGvRg		241
srnX		dyltLFsRae	YEaAYrkAGl	dVeYvvtgeg	spGFFlGtRr		
rdmD		hiisLFsRse	YEqaFrdAGf	aVeYlpeaps	grGLFvGvRg		
Consensus		-----LF-R--	YE-AF--AG-	-V-Y-----	--GLF-G-R-	-	

FIG. 5E-2

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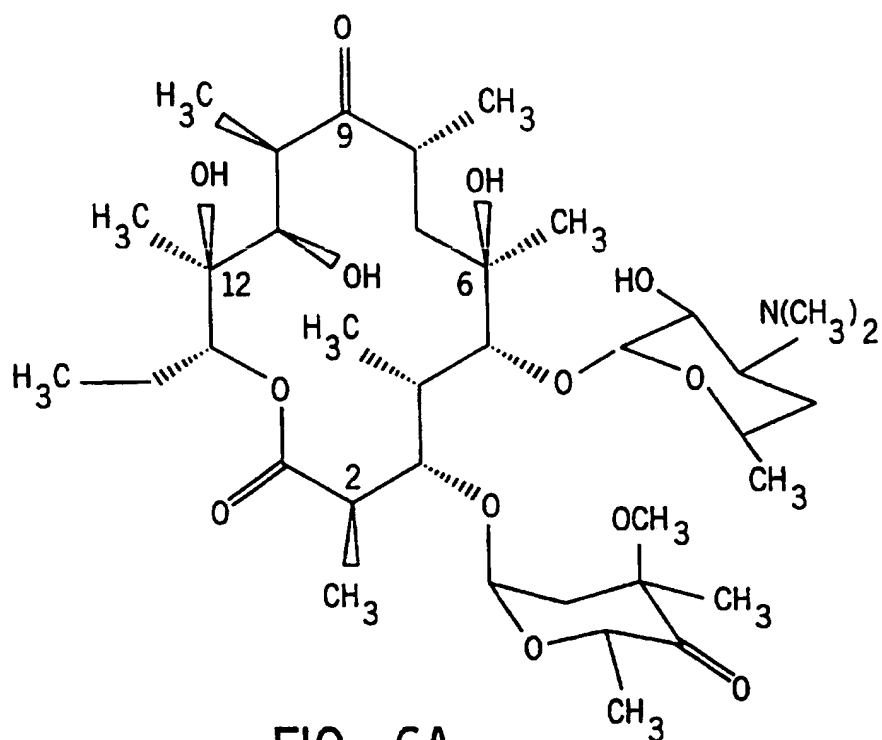


FIG. 6A

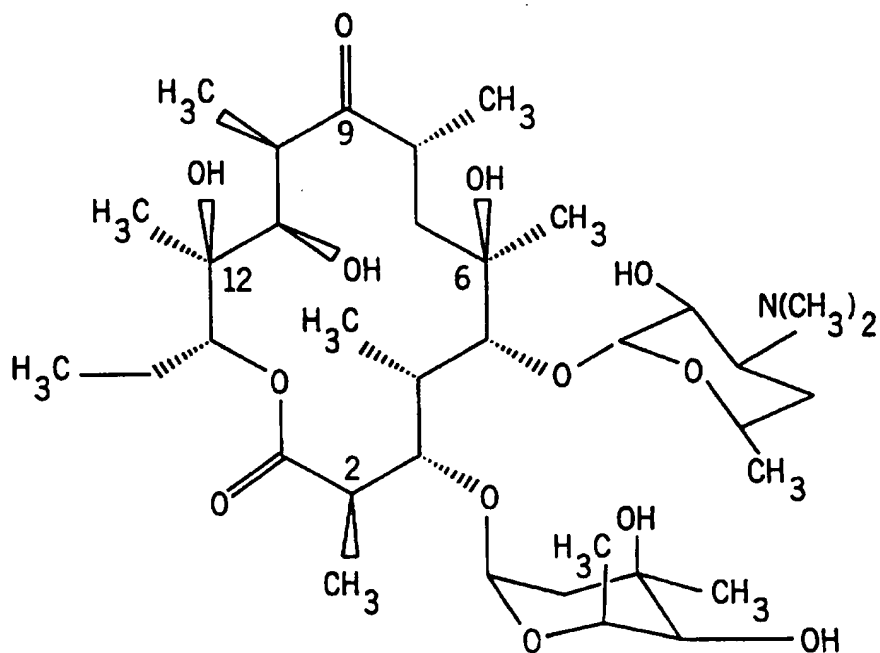


FIG. 6B

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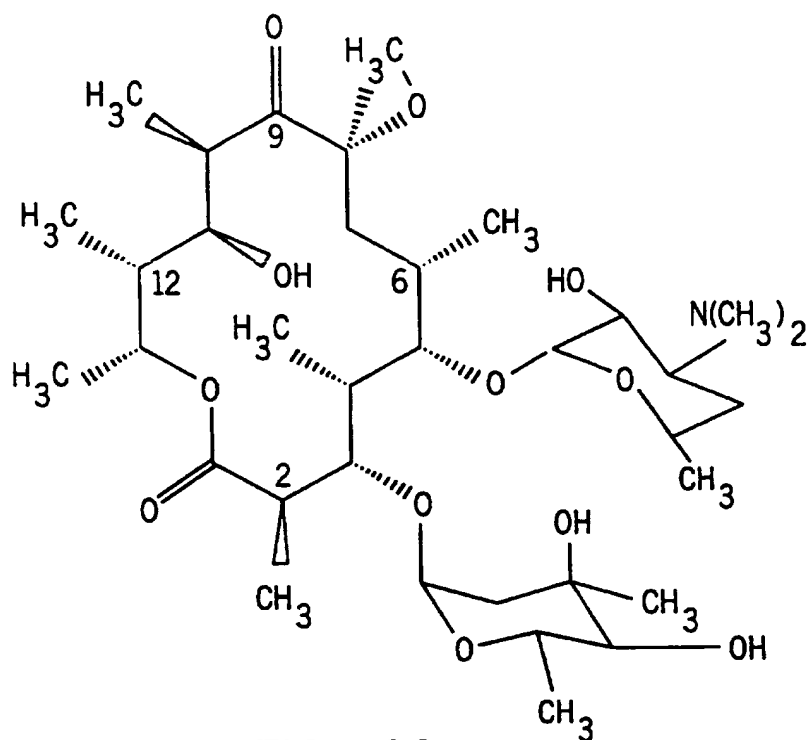


FIG. 6C

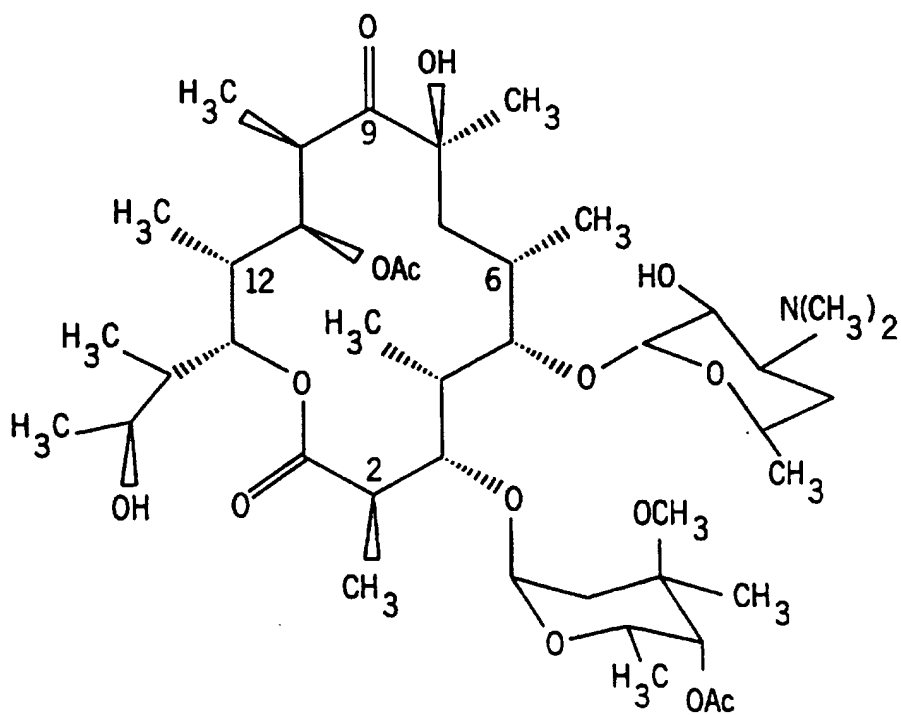


FIG. 6D

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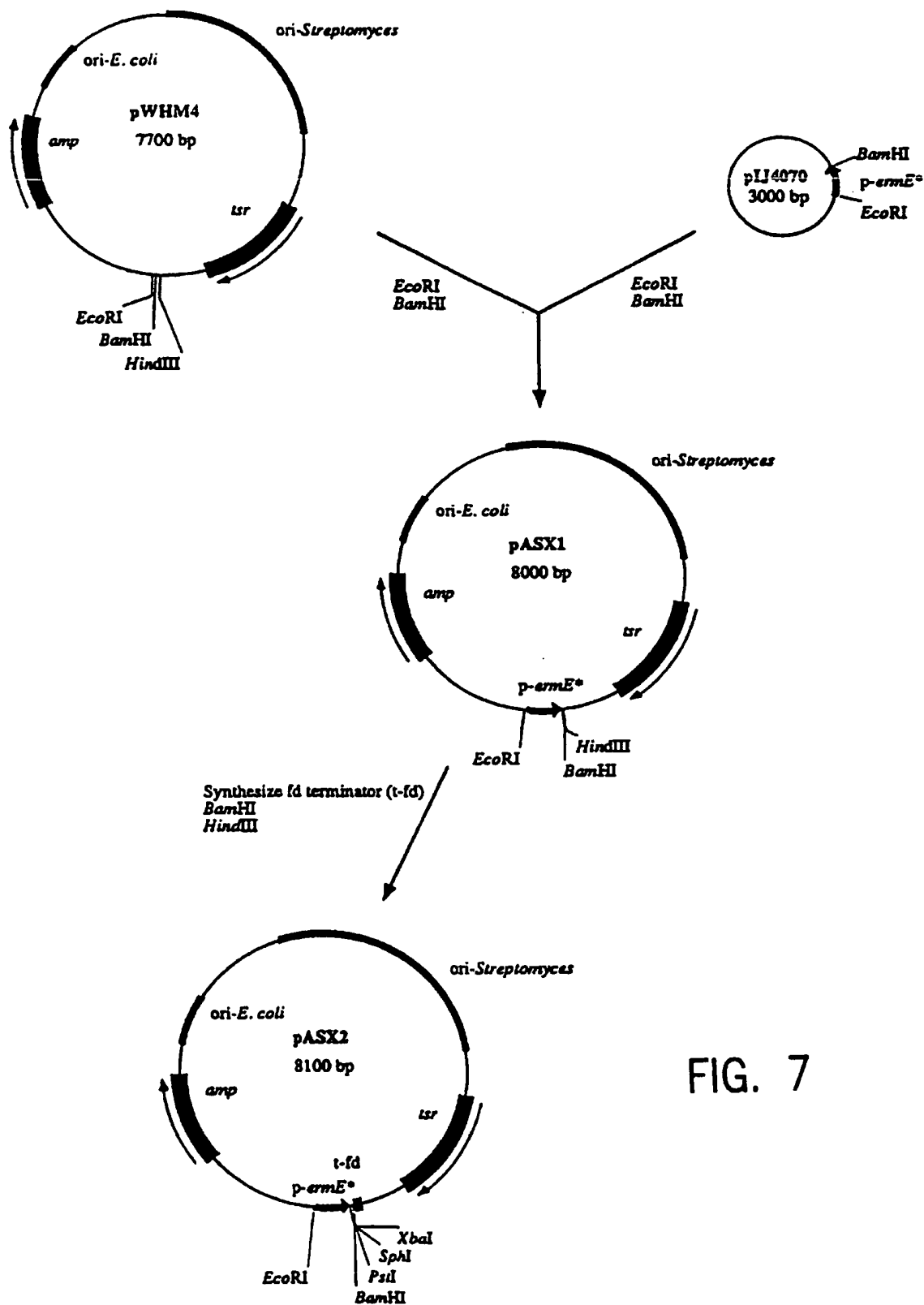


FIG. 7

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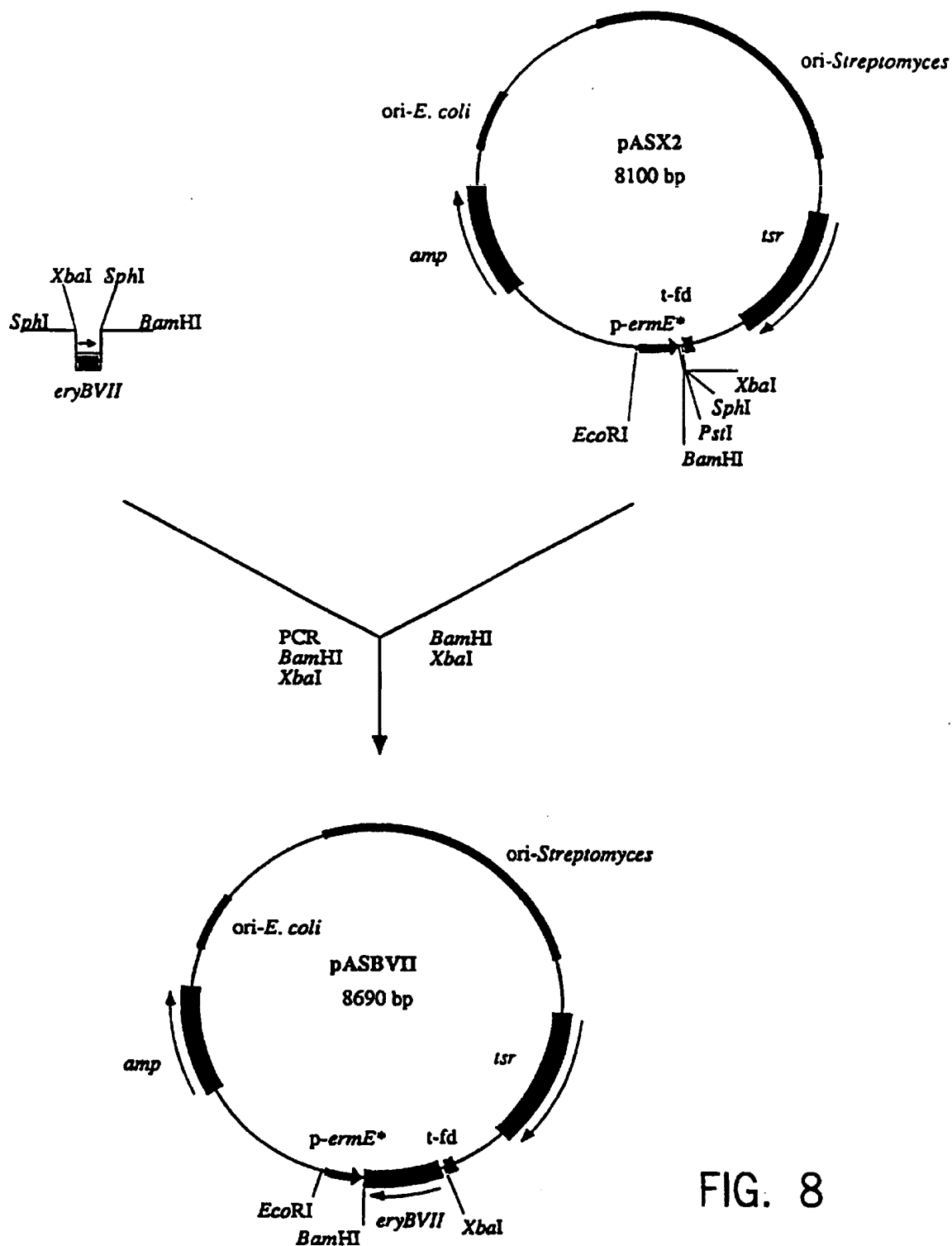


FIG. 8

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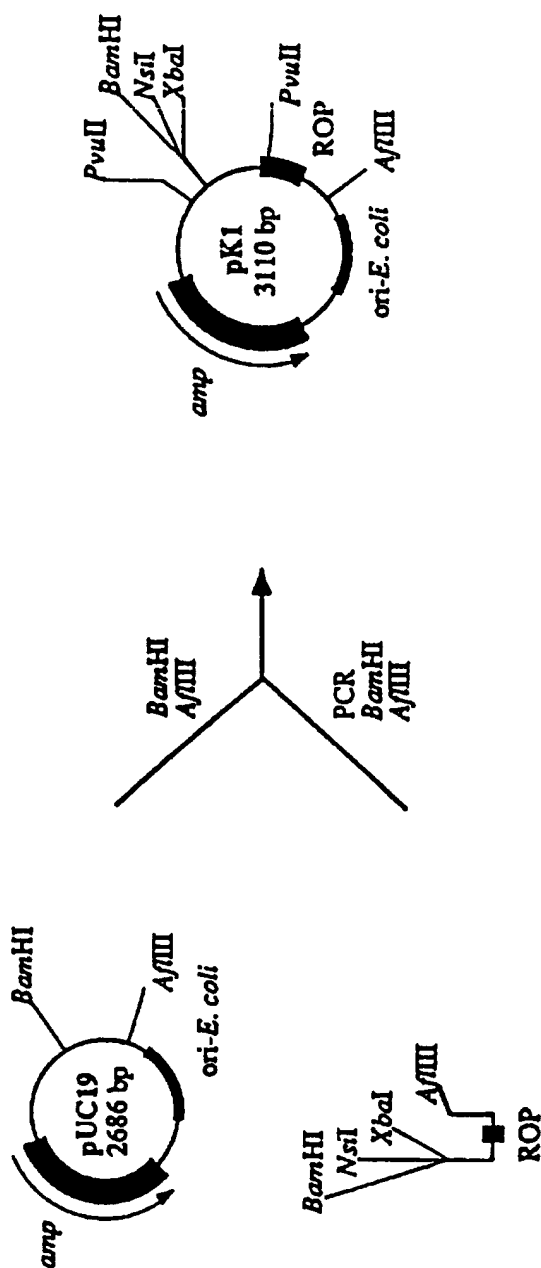


FIG. 9A

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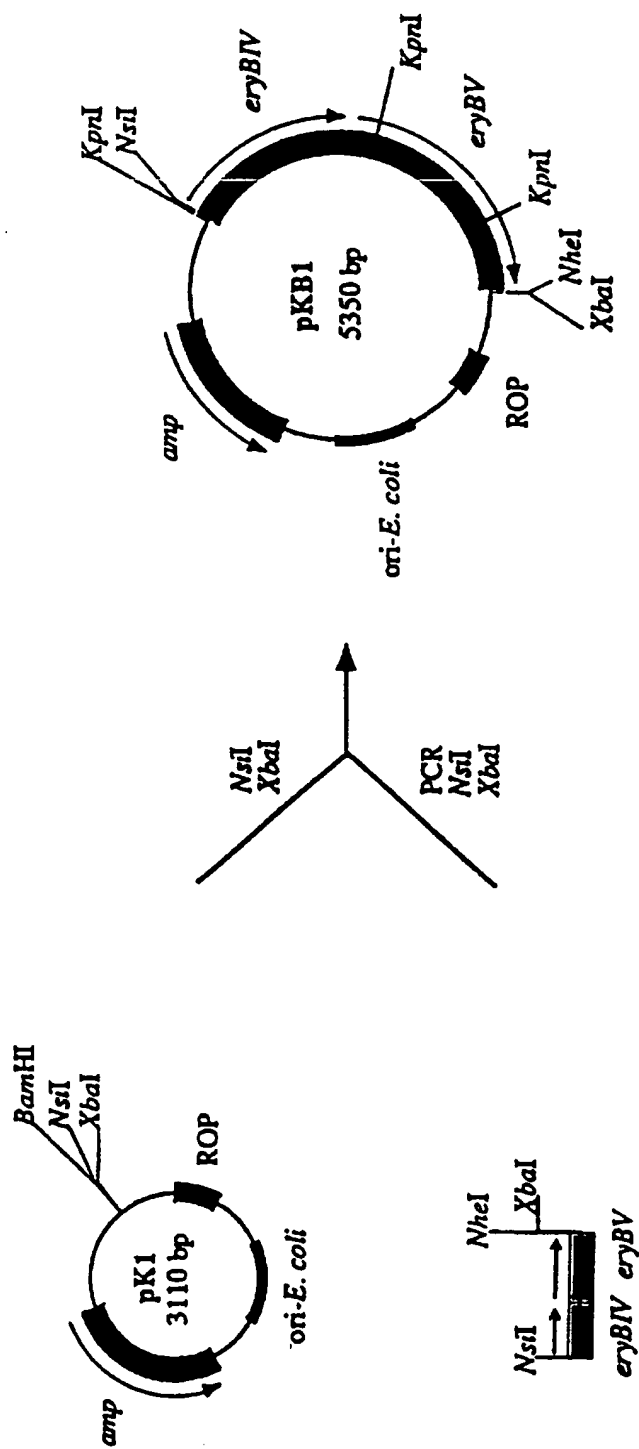


FIG. 9B

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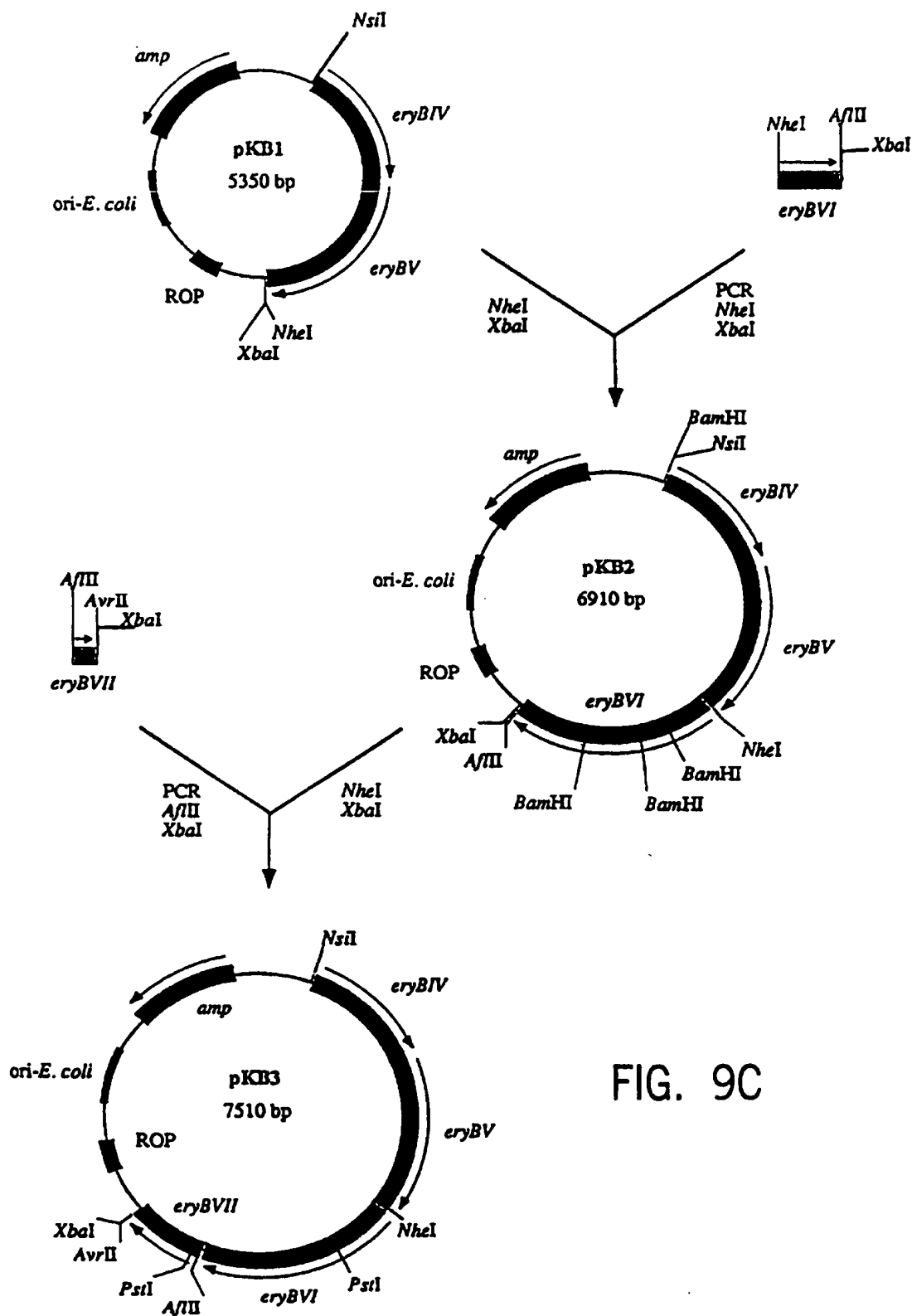


FIG. 9C

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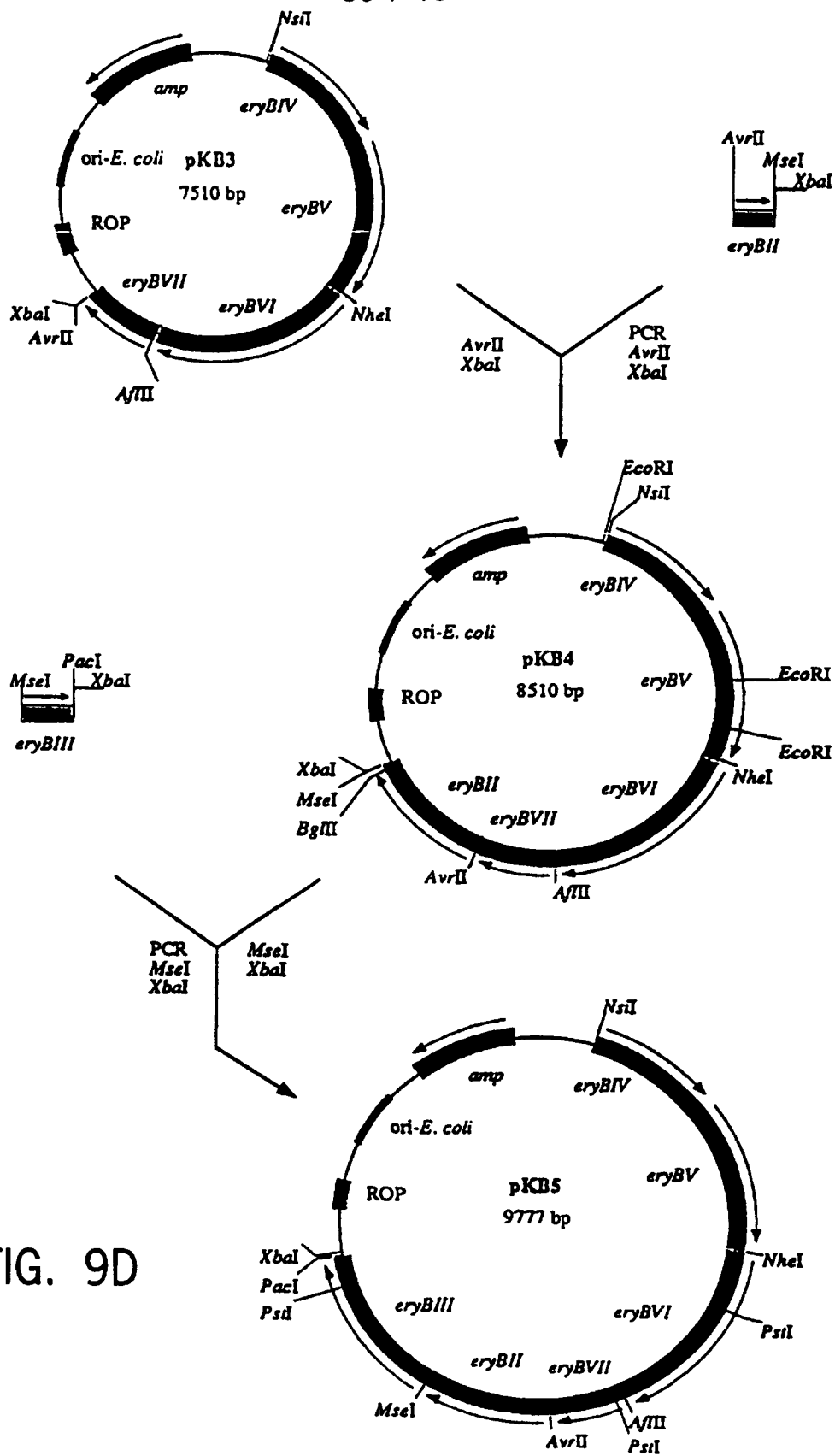


FIG. 9D

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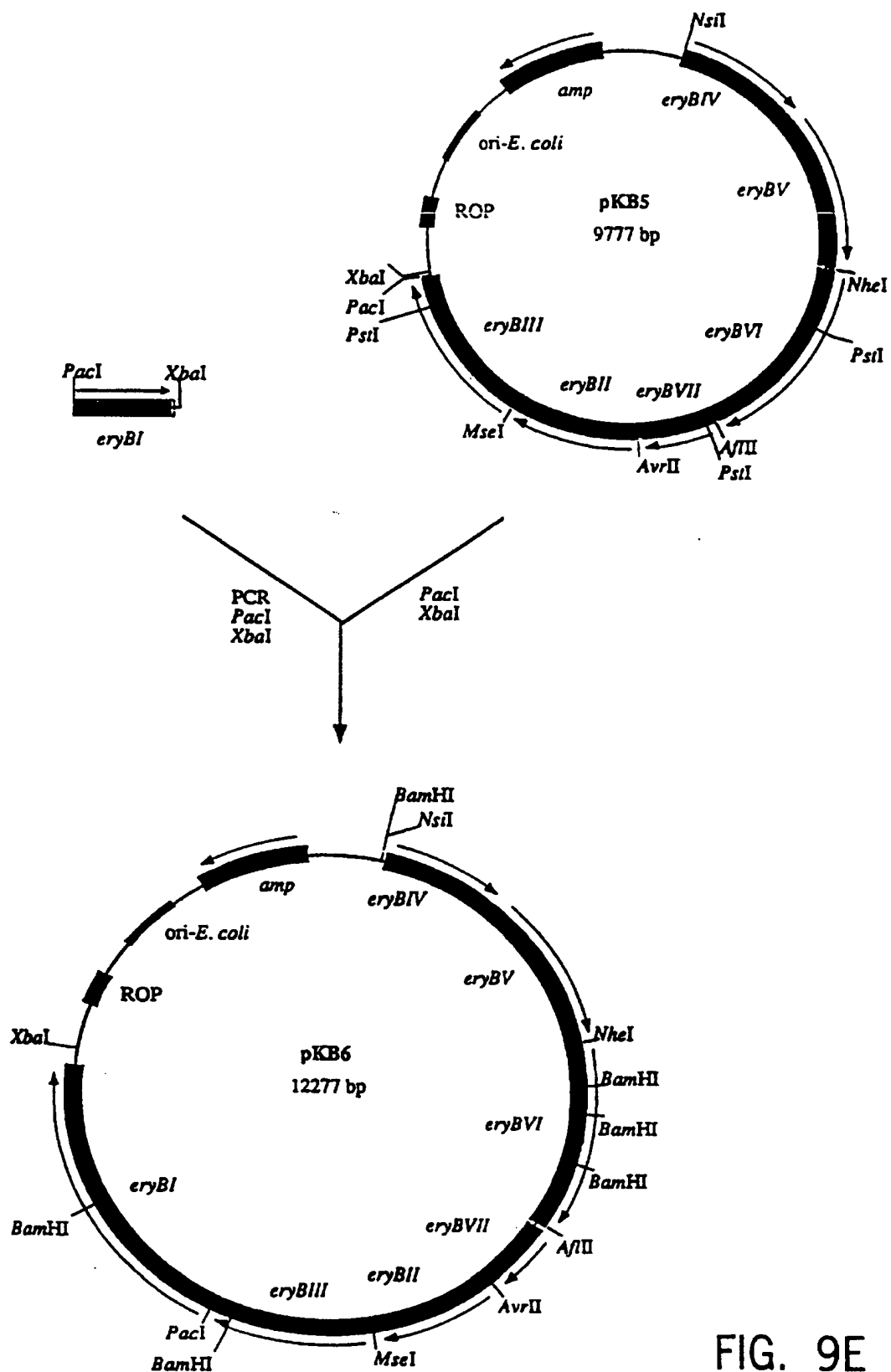


FIG. 9E

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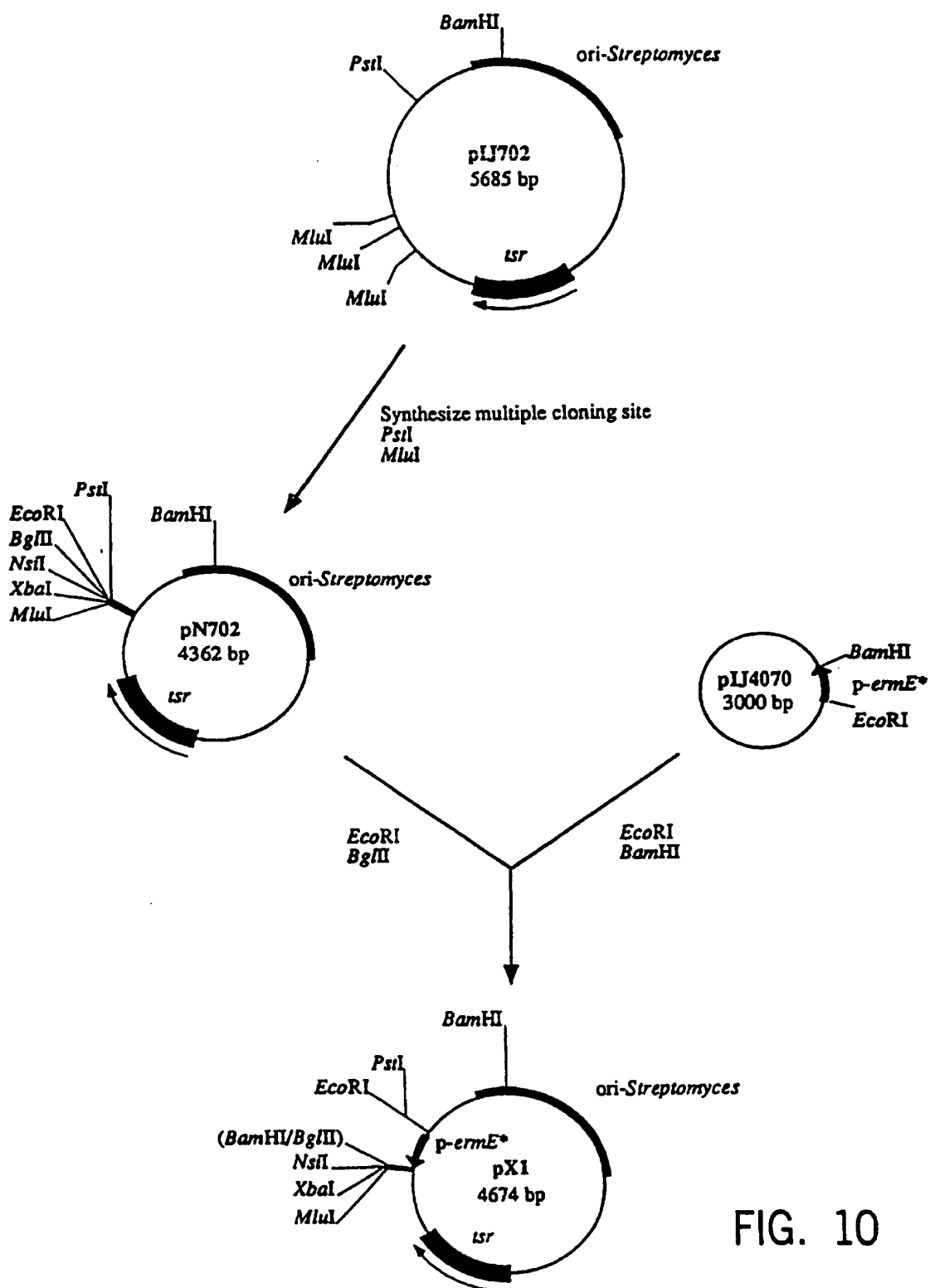


FIG. 10

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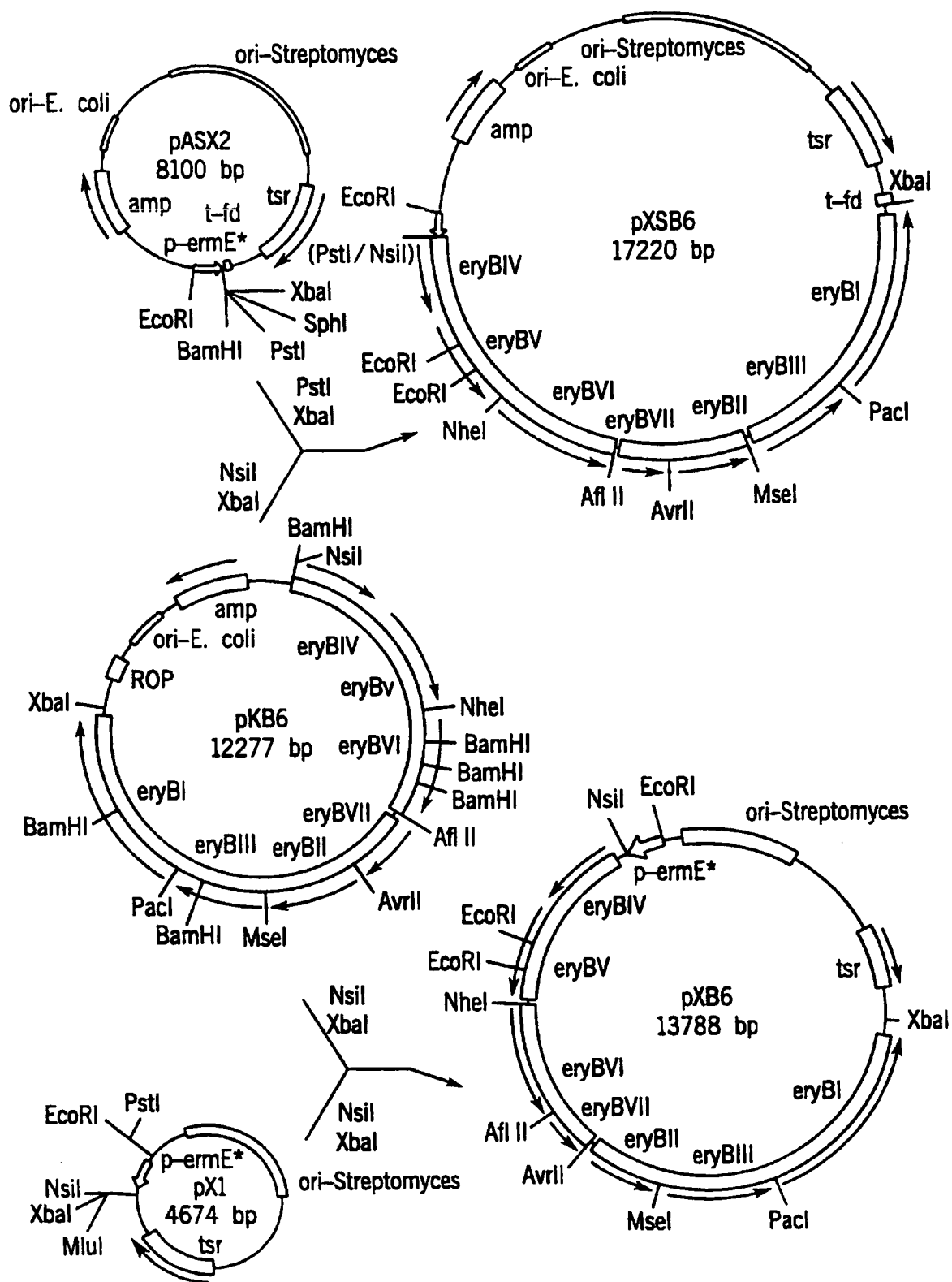


FIG. 11

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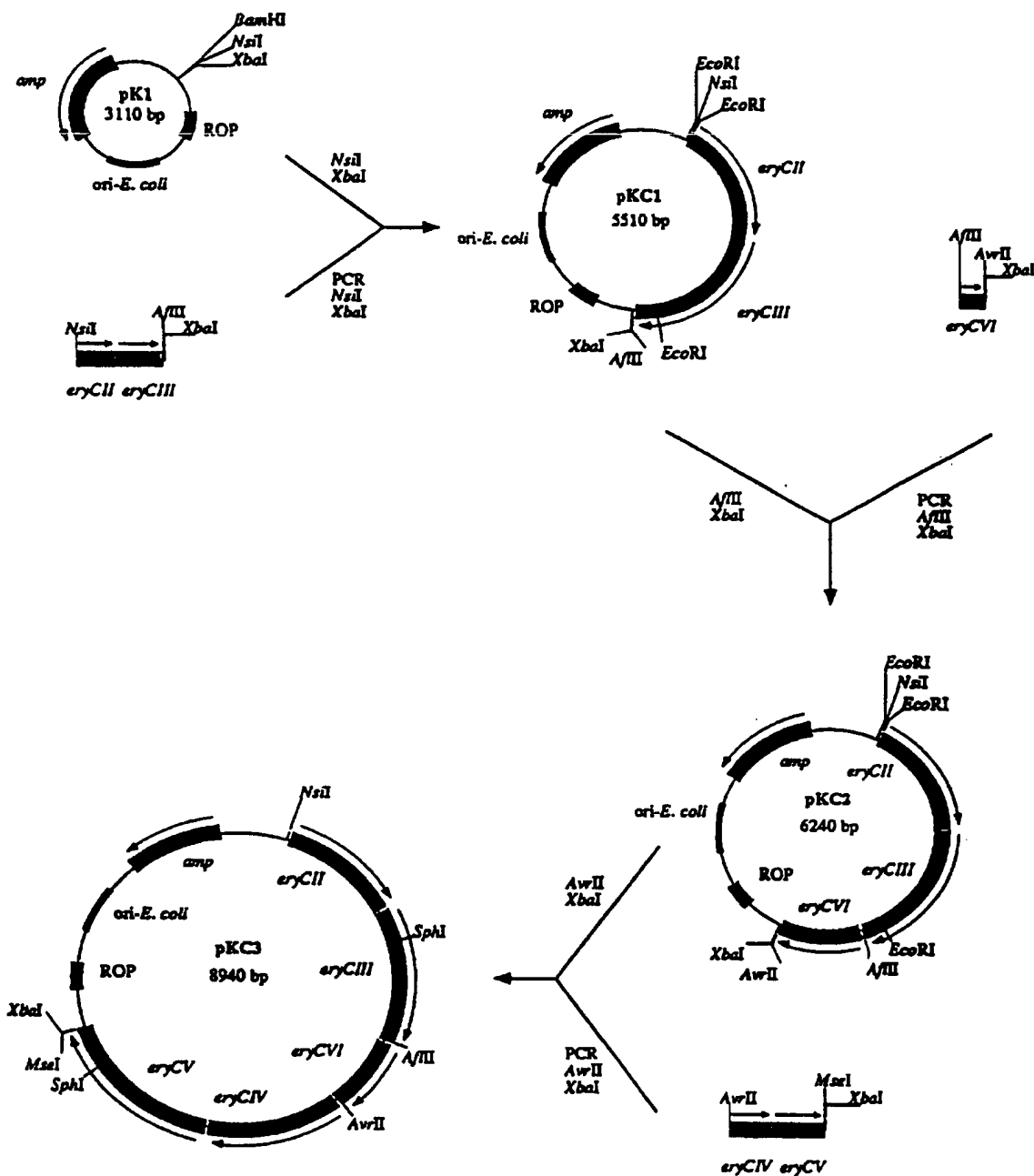


FIG. 12A

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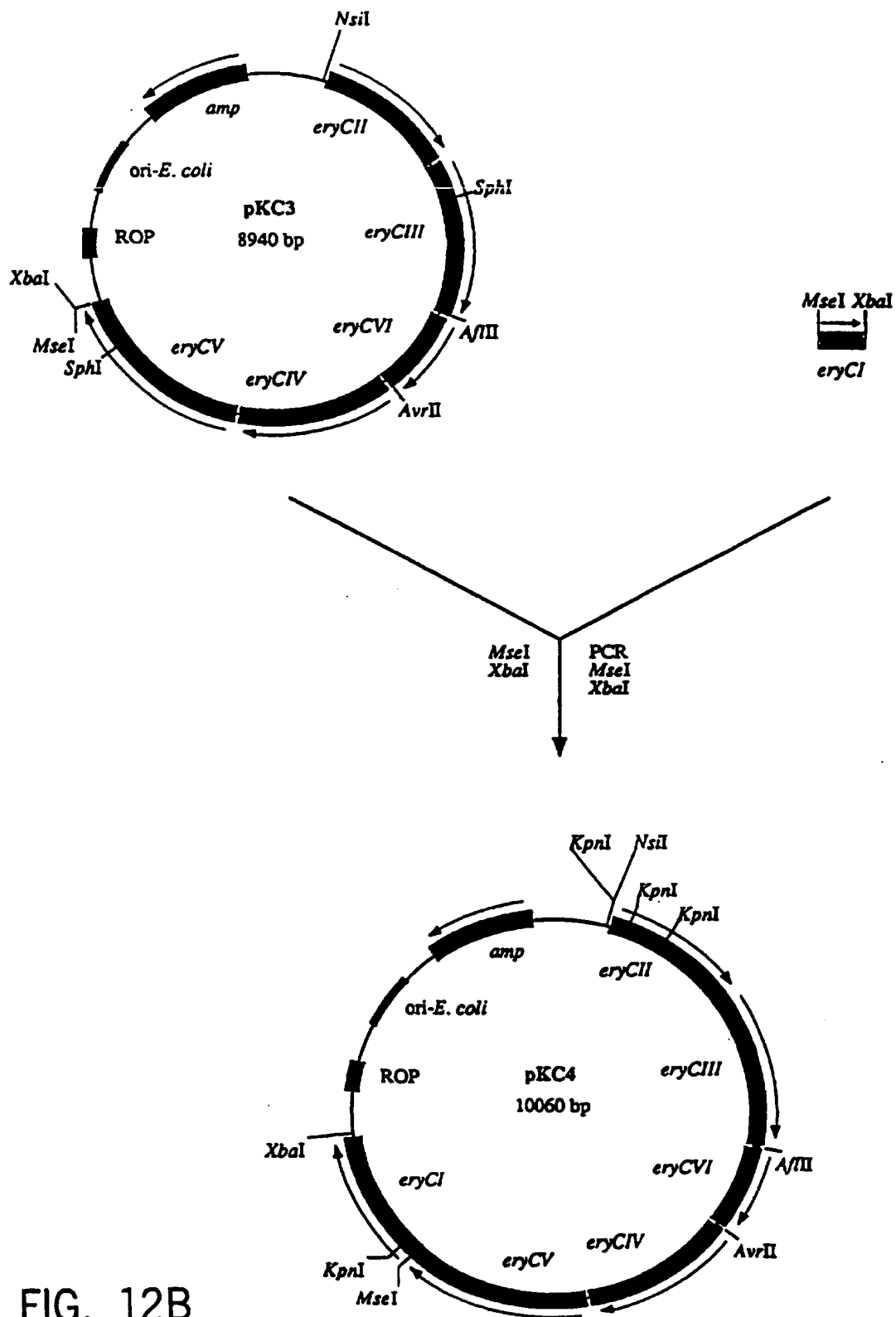


FIG. 12B

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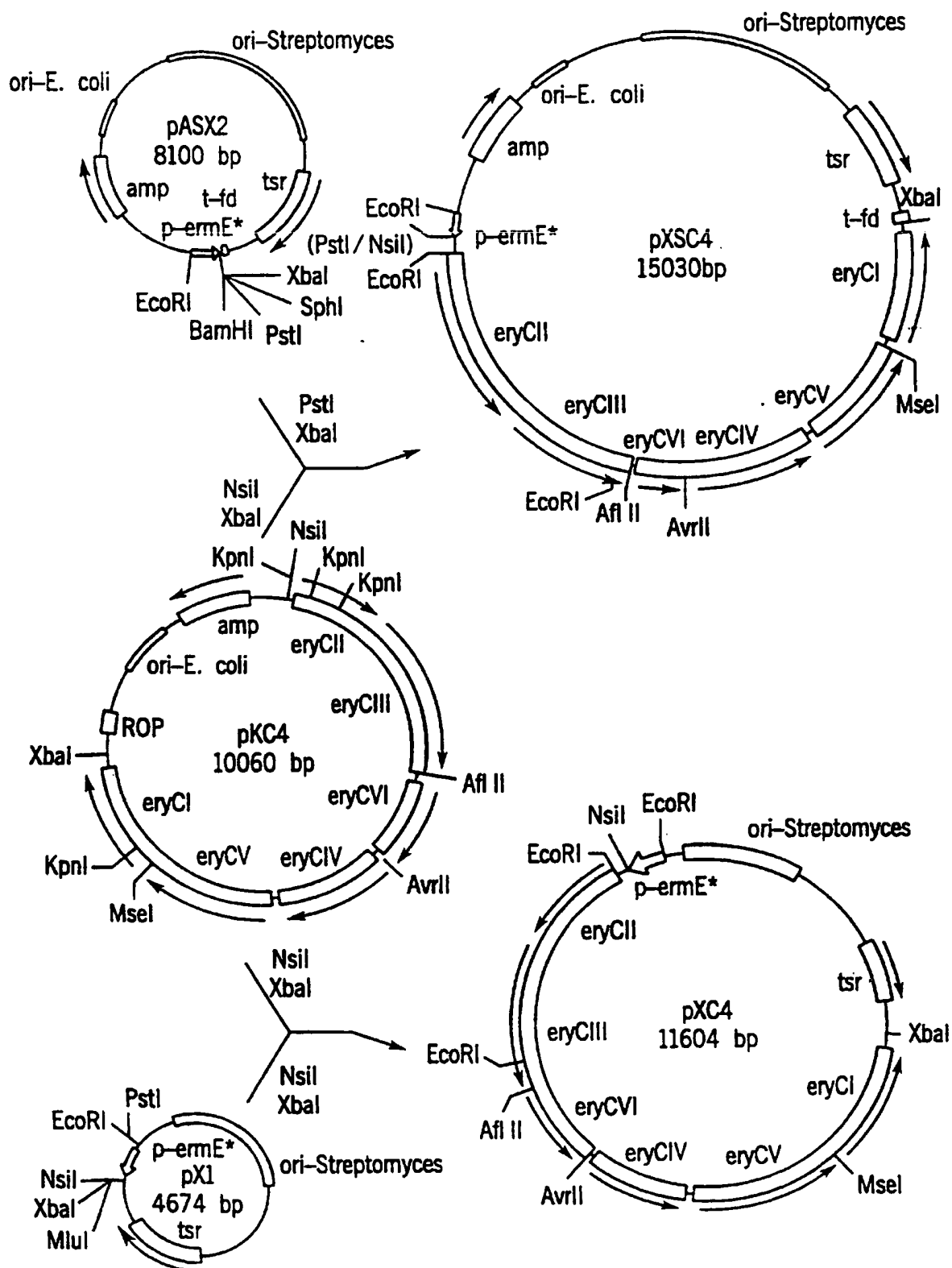


FIG. 13

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